(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau

(43) International Publication Date
13 June 2002 (13.06.2002)

PCT

(10) International Publication Number
WO 02/46228 A2

(51) International Patent Classification[7]: C07K 14/705

(21) International Application Number: PCT/US01/30941

(22) International Filing Date: 3 October 2001 (03.10.2001)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
60/251,481 5 December 2000 (05.12.2000) US

(71) Applicant: **WISCONSIN ALUMNI RESEARCH FOUNDATION** [US/US]; 614 Walnut Street, P.O. Box 7365, Madison, WI 53707-7365 (US).

(72) Inventors: **YOUNG, John, A., T.**; 1154 Sherman Avenue, Madison, WI 53706 (US). **BRADLEY, Kenneth, A.**; 616 North Carrol Street #305, Madison, WI 53703 (US). **COLLIER, Robert, J.**; 43 Garden Road, Wellesley, MA 02481 (US). **MOGRIDGE, Jeremy, S.**; 7 Atherton Road, Apt. 2, Brookline, MA 02446 (US).

(74) Agent: **BERSON, Bennett, J.**; Quarles & Brady LLP, P.O. Box 2113, Madison, WI 53701-2113 (US).

(81) Designated States *(national)*: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW.

(84) Designated States *(regional)*: ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:
— *without international search report and to be republished upon receipt of that report*

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

WO 02/46228 A2

(54) Title: RECEPTOR FOR *B. ANTHRACIS* TOXIN

(57) Abstract: The present invention relates to mammalian anthrax toxin receptor polypeptides and polynucleotides encoding same as well as related polypeptides and polynucleotides, vectors containing the polynucleotides and polypeptides, host cells containing related polynucleotide molecules, and cells displaying no anthrax toxin receptor on an exterior surface of the cells-minus cell lines and animals. The present invention also relates to methods for identifying molecules that bind the anthrax toxin receptor and molecules that reduce the toxicity of anthrax toxin. Finally, the present invention provides methods for treating human and non-human animals suffering from anthrax.

RECEPTOR FOR B. ANTHRACIS TOXIN

CROSS-REFERENCE TO RELATED APPLICATIONS

[0001] This application claims the benefit of U.S. provisional application serial number 60/251,481, filed on December 5, 2000, which is incorporated herein by reference as if set forth in its entirety.

STATEMENT REGARDING FEDERALLY SPONSORED RESEARCH OR DEVELOPMENT

[0002] Not applicable.

BACKGROUND OF THE INVENTION

[0003] *Bacillus anthracis*, the spore-forming causative agent of anthrax, generally infects herbivores (Hanna, 1998). Human infection, while rare, can result in a generally benign, self-limiting cutaneous disease or a systemic disease that rapidly leads to death in a high percentage of cases. The cutaneous disease can arise when spore particles from soil or animal products are introduced into cuts or skin abrasions. In contrast, the systemic disease can arise when *B. anthracis* spore particles are inhaled ($LD_{50} \approx$ 10,000 spore particles). The high mortality rate and the ability to readily prepare and deliver *B. anthracis* spore particles as an aerosol have made *B. anthracis* a dreaded agent of biowarfare and bioterrorism.

[0004] The causative agent of the systemic disease is anthrax toxin (AT), which itself comprises a pair of binary, AB-type toxins -- lethal toxin and edema toxin (Leppla, 1995). Each is assembled at the surface of mammalian cells from proteins released by *B. anthracis*. Lethal toxin, assembled from Protective Antigen (PA, 83kDa) and Lethal Factor (LF, 90 kDa), is primarily responsible for lethality (Friedlander, 1986; Hanna et al., 1992; Hanna et al., 1993). Edema toxin, assembled from PA and Edema Factor (EF, 89 kDa), causes edema at the site of injection (Leppla, 1982). EF has calmodulin-dependent adenylate cyclase activity. LF is a Zn^{++}-dependent protease that cleaves certain proteins involved in signal transduction and cell cycle progression (MAPKK1 and MAPKK2) (Duesbery et al., 1998).

[0005] In these AB-type toxins, PA is the receptor-binding B moiety that delivers either EF or LF, as alternative enzymic A moieties, to the cytosol of mammalian cells (Leppla, 1995). Initially, PA binds specifically, reversibly, and with high affinity (Kd ≈ 1 nM) to a cell-surface AT receptor (ATR). After binding to the receptor, PA is cleaved by a member of the furin family of proprotein convertases, which removes a 20 kDa fragment, PA20, from the N-terminus (Klimpel et al., 1992; Novak et al., 1992). The complementary fragment, PA63, remains receptor-bound and spontaneously self-associates to form heptameric ring-shaped oligomers (Milne et al., 1994) that avidly and competitively bind EF and/or LF (Leppla, 1995) to form EF/LF-PA63 complexes. These complexes are trafficked to an acidic compartment by receptor-mediated endocytosis. In the acidic compartment, the PA63 heptamers (the "prepore") are inserted into the membrane, forming transmembrane pores (Gordon et al., 1988). Concomitantly EF and LF are translocated across the membrane to the cytosol. Consistent with the pH dependence of translocation, toxin action is inhibited by lysosomotropic agents and bafilomycin A1 (Mendard et al., 1996).

[0006] EF translocation causes a large increase in intracellular cAMP concentration (Gordon et al., 1988; Gordon et al., 1989). Increased cAMP levels cause edema, and in neutrophils, inhibit phagocytosis and oxidative burst (O'Brien et al., 1985). By protecting the bacteria from phagocytosis, edema toxin apparently aids in establishing bacterial infection and proliferation in the mammalian host.

[0007] Treatment of primary macrophages and certain macrophage cell lines with lethal toxin causes cell lysis (Friedlander, 1986). Macrophage-depleted mice are resistant to treatment with lethal toxin, suggesting that macrophages are the primary targets of lethal toxin (Hanna et al., 1993). Low doses of lethal toxin induce the production of interleukin-1 and tumor necrosis factor (Hanna et al., 1993). Thus, it has been suggested that hyperproduction of cytokines causes death of the host by inducing systemic shock. How these or other proteins lead to cytokine production and macrophage lysis remains unclear.

[0008] In the past few years considerable progress has been made toward a detailed understanding of the structure and function of PA. Crystallographic structures of PA and the PA63 heptamers have been determined (Petosa et al., 1997). The prepore undergoes a major conformational change under acidic conditions to form a 14-strand transmembrane β-barrel pore

(Benson et al., 1998; Miller et al., 1999). The pore structure and the detailed mechanism by which LF and EF are translocated across membranes are under intensive investigation.

[0009] The ATR structure is heretofore unknown, but is present in all cell lines that have been tested. Studies on CHO-K1 cells had indicated that PA binds to a proteinaceous receptor that is present in about 10^4 copies/cell (Escuyer and Collier, 1991). The paucity of knowledge about the ATR represents a major gap in the understanding of how AT acts. Identification and cloning of the ATR will provide more treatment strategies for anthrax.

[00010] A cDNA clone (Genbank Accession Number NM 032208) known as tumor endothelial marker 8 (TEM8) is known (St. Croix, 2000). TEM8 is upregulated in colorectal cancer endothelium, but heretofore the function of TEM8 was not known.

BRIEF SUMMARY OF THE INVENTION

[00011] The present application discloses structures of complete and partial anthrax toxin receptors from a mammal, namely a human. The complete anthrax toxin receptor includes an extracellular domain, a transmembrane domain, and a cytoplasmic domain that can vary in length, as is disclosed herein. It is disclosed herein that PA binds to the anthrax toxin receptor at a von Willebrand factor A (VWA) domain in the extracellular domain.

[00012] In one aspect, the invention is summarized in that an anthrax toxin receptor is a polypeptide having an amino acid sequence selected from SEQ ID NO:2, SEQ ID NO:6, SEQ ID NO: 8, SEQ ID NO:10, a PA-binding fragment of any of the foregoing, and a PA-binding variant of any of the foregoing polypeptides having conservative or non-conservative amino acid substitutions or other changes relative to the disclosed sequences. The various forms of the receptor encoded by SEQ ID NO:2, SEQ ID NO:6, SEQ ID NO: 8, and SEQ ID NO:10 apparently differ as a result of alternative splicing.

[00013] In a related aspect, the invention further relates to an isolated polynucleotide that encodes any of the above-mentioned polypeptides and their complements, and a polynucleotide that hybridizes under moderately stringent or stringent hybridization conditions to any of the foregoing.

[00014] In still another related aspect, the invention encompasses a cloning vector and an expression vector comprising any of the foregoing polynucleotides, whether or not the

polynucleotide is operably linked to an expression control sequence that does not natively promote transcription or translation of the polynucleotide.

By identifying the polypeptides and polynucleotides of the invention, the applicant enables the skilled artisan to detect and quantify mRNA and ATR protein in a sample, and to generate *atr* transgenic and *atr* knock-out animals using methods available to the art.

[00015] Further, the invention includes a host cell comprising any such vector in its interior. Also within the scope of the present invention is a host cell having a polynucleotide of the invention integrated into the host cell genome at a location that is not the native location of the polynucleotide.

[00016] In yet another aspect, the invention is a method for producing an anthrax toxin receptor polypeptide that includes the steps of transcribing a polynucleotide that encodes an anthrax toxin receptor polypeptide, operably linked to an upstream expression control sequence, to produce an mRNA for the receptor polypeptide, and translating the mRNA to produce the receptor polypeptide. This method can be performed in a host cell when the polynucleotide is operably linked to the expression control sequence in an expression vector, and wherein the expression vector is delivered into a host cell, the expression control sequence being operable in the host cell. Alternatively, at least one of the transcribing and translating steps can be performed in an *in vitro* system, examples of which are well known in the art and commercially available. In either case, the polypeptide can be isolated from other cellular material using readily available methods.

[00017] In still another aspect, the invention is a method for identifying an agent that can alter the effect of AT on the host cell or organism. The method includes the steps of separately exposing a plurality of putative agents in the presence of AT to a plurality of cells having on their surface at least a portion of the ATR that binds to AT or a component thereof, comparing the effect of AT on the cells in the presence and absence of the agent, and identifying at least one agent that alters an effect of AT on the cells. In a related aspect, the present invention encompasses an agent that alters binding of AT to the ATR.

[00018] The present invention also encompasses a method for reducing or preventing AT-related damage *in vivo* or *in vitro* to human or non-human cells having an ATR on an outer cell

surface, the method comprising the step of exposing the cells to an agent that reduces binding of AT to the ATR. Similarly, the invention relates to a method for reducing or preventing damage *in vivo* or *in vitro* to human or non-human cells caused by AT by exposing AT to an agent that reduces binding of the AT to the ATR.

[00019] The present invention is also a method for identifying a mutant of the extracellular ATR domain or fragment thereof having altered (increased or reduced) binding affinity for AT.

[00020] It is an object of the invention to identify polypeptides that encode a mammalian anthrax toxin receptor, as well as fragments, mutants, and variants thereof and polynucleotides encoding same.

[00021] It is a feature of the invention that a soluble PA-binding polypeptide can reduce or eliminate toxicity associated with anthrax toxin.

[00022] Other objects, advantages and features of the invention will become apparent from the following specifications and claims.

BRIEF DESCRIPTION OF THE SEVERAL VIEWS OF THE DRAWINGS

[00023] Fig. 1 shows sequence alignment of various ATR polypeptide sequences with the I domain of integrin α2 and with the von Willebrand factor A domain consensus sequence.

DETAILED DESCRIPTION OF THE INVENTION

[00024] An isolated polynucleotide and an isolated polypeptide, as used herein, can be isolated from its natural environment or can be synthesized. Complete purification is not required in either case. Amino acid and nucleotide sequences flanking an isolated polypeptide or polynucleotide that occurs in nature, respectively, can but need not be absent from the isolated form.

[00025] Further, an isolated polynucleotide has a structure that is not identical to that of any naturally occurring nucleic acid or to that of any fragment of a naturally occurring genomic nucleic acid spanning more than three separate genes. The term includes, without limitation, (a) a nucleic acid molecule having a sequence of a naturally occurring genomic or extrachromosomal nucleic acid molecule but which is not flanked by the coding sequences that flank the sequence in its natural position; (b) a nucleic acid molecule incorporated into a vector or into a prokaryote or

eukaryote genome such that the resulting molecule is not identical to any naturally occurring vector or genomic DNA; (c) a separate molecule such as a cDNA, a genomic fragment, a fragment produced by polymerase chain reaction (PCR), or a restriction fragment; and (d) a recombinant nucleotide sequence that is part of a hybrid gene, i.e., a gene encoding a fusion protein. Specifically excluded from this definition are nucleic acids present in mixtures of clones, e.g., as these occur in a DNA library such as a cDNA or genomic DNA library. An isolated nucleic acid molecule can be modified or unmodified DNA or RNA, whether fully or partially single-stranded or double-stranded or even triple-stranded. A nucleic acid molecule can be chemically or enzymatically modified and can include so-called non-standard bases such as inosine.

[00026] Reference herein to use of AT is understood to encompass use of an ATR-binding component thereof, especially PA.

Anthrax Toxin Receptor

[00027] The applicants have identified and determined the nucleic acid sequence (SEQ ID NO:1) of a cDNA clone that of a 368 amino acid long polypeptide (SEQ ID NO:2, *ATR*), and show herein that the polypeptide is a surface-bound anthrax toxin receptor (ATR) on human cells. Based on known structural analysis methods, the polypeptide is predicted to encode a 27 amino-acid-long signal peptide (amino acids 1-27 of SEQ ID NO:2), a 293 amino-acid-long extracellular domain (amino acids 28-320 of SEQ ID NO:2), a 23 amino-acid-long putative transmembrane region (amino acids 320-343 of SEQ ID NO:2), and a 25 amino acid long cytoplasmic domain (amino acids 344-368 of SEQ ID NO:2).

[00028] It is disclosed herein that Protective Antigen (PA) of anthrax toxin (AT) binds to the anthrax toxin receptor at a von Willebrand factor A (VWA) domain located in the portion from amino acid 44 to 216 in the extracellular domain of SEQ ID NO:2. VWA domains are present in the extracellular portions of a variety of cell surface proteins, including matrilins and integrins (designated as I domains). A VWA domain consensus sequence, VWA-CON, developed by comparing 210 related sequences, is presented as SEQ ID NO:3. These domains are important for protein/protein interactions and constitute ligand binding sites for integrins (Dickeson, 1998). The I domain of integrin α2 (α2) is presented as SEQ ID NO:4. Ligand binding through I domains requires an intact metal ion-dependent adhesion site (MIDAS) motif (Lee,

1995) which appears to be conserved in the ATR extracellular domain, as is detailed below.

[00029] Comparison of SEQ ID NO:1 and SEQ ID NO:2 to existing databases revealed other versions of those sequences. Human cDNA TEM8 (SEQ ID NO:5; Genbank accession number NM 032208) encodes a 564 amino-acid-long form (SEQ ID NO:6) of the human ATR. SEQ ID NO:6 has not previously been identified as an anthrax toxin receptor, and indeed no function has yet been ascribed to the protein. Like SEQ ID NO:1, SEQ ID NO:5 was a PCR amplification product from HeLa cells and human placenta cDNA libraries. Whereas the cytoplasmic tail of SEQ ID NO:2 is only 25 amino acids long, that of SEQ ID NO:6 is predicted to be 221 amino acids long (amino acids 344-564), presumably as a result of differential splicing of a primary mRNA transcript. The proteins are otherwise identical. Upstream of the coding sequences, SEQ ID NO:1 and SEQ ID NO:5 are also identical.

[00030] Also presented are IMAGE CLONE 4563020 (SEQ ID NO:7; Genbank Accession Number BC012074) and the predicted polypeptide encoded by the clone (SEQ ID NO:8). SEQ ID NO:8 is identical to amino acids 1-317 of ATR, but differs thereafter at the C-terminus. Similarly, human cDNA FLJ10601, clone NT2RP2005000 (SEQ ID NO:9; Genbank Accession Number AK001463) and the predicted polypeptide encoded by the clone (SEQ ID NO:10) are presented. This polypeptide is identical to a portion of SEQ ID NO:2 from amino acid 80 to amino acid 218. As with TEM8 and the protein it encodes, no function is known for any of these polynucleotide and polypeptide sequences, nor has there been any prior indication that the polypeptides are complete or partial anthrax toxin receptors.

[00031] It is of interest to note that the product of the mouse homolog of ATR/TEM8 (Genbank accession number AK013005) is highly related to the human clones, sharing greater than 98% amino acid sequence identity within the reported extracellular domain. This suggests that the anthrax toxin receptor is conserved among species. Furthermore, consistent with the observation that the anthrax toxin receptor is found in a variety of cell lines, *ATR* is expressed in a number of different tissues including CNS, heart, lung, and lymphocytes.

[00032] In addition to the full-length and partial ATR polypeptide sequences presented in SEQ ID NO:2, SEQ ID NO:6, SEQ ID NO:8 and SEQ ID NO:10, other polypeptide fragments shorter than those sequences that retain PA-binding activity, and variants thereof are also within the scope of the invention. The entire receptor is not required for utility; rather, fragments that

bind to PA are useful in the invention.

[00033] A skilled artisan can readily assess whether a fragment binds to PA. A polypeptide is considered to bind to PA if the equilibrium dissociation constant of the binary complex is 10 micromolar or less. PA-binding to the ATR (or a fragment of the ATR) can be measured using a protein-protein binding method such as coimmunoprecipitation, affinity column analysis, ELISA analysis, flow cytometry or fluorescence resonance energy transfer (FRET), and surface plasmon resonance (SPR). SPR is particularly suited as it is highly sensitive and accurate, operable in real time, and consumes only minute amounts of protein. SPR uses changes in refractive index to quantify macromolecular binding and dissociation to a ligand covalently tethered to a thin gold chip in a micro flow cell. Besides the equilibrium dissociation constant (Kd), on- and off-rate constants (ka and kd) can also be obtained. A BIAcore 2000 instrument (Pharmacia Biotech) can be used for these measurements. Typically, a protein is covalently tethered to a carboxymethyl dextran matrix bonded to the gold chip. Binding of a proteinaceous ligand to the immobilized protein results in a quantifiable change in refractive index of the dextran/protein layer. SPR can also be used to determine whether the interaction between PA and its receptor is sensitive to low pH, which is relevant to toxin endocytosis. This technique has been used to study protein-protein interactions in many systems, including the interactions of PA63 with EF and LF (Elliott, 1998).

[00034] The invention also relates to polypeptides that are at least 80%, preferably at least 90%, more preferably at least 95%, still more preferably at least 97%, or most preferably at least 99% identical to any aforementioned PA-binding polypeptide fragment, where PA-binding is maintained. As used herein, "percent identity" between amino acid or nucleic acid sequences is synonymous with "percent homology," which can be determined using the algorithm of Karlin and Altschul (Proc. Natl. Acad. Sci. USA 87:2264-2268, 1990), modified by Karlin and Altschul (Proc. Natl. Acad. Sci. USA 90:5873-5877, 1993). Such an algorithm is incorporated into the NBLAST and XBLAST programs of Altschul et al. (J. Mol. Biol. 215:403-410, 1990). BLAST nucleotide searches are performed with the NBLAST program, score = 100, wordlength = 12, to obtain nucleotide sequences homologous to a nucleic acid molecule of the invention. BLAST protein searches are performed with the XBLAST program, score = 50, wordlength = 3, to obtain amino acid sequences homologous to a reference polypeptide (e.g., SEQ ID NO:2). To obtain

gapped alignments for comparison purposes, Gapped BLAST is utilized as described in Altschul et al. (Nucleic Acids Res. 25:3389-3402, 1997). When utilizing BLAST and Gapped BLAST programs, the default parameters of the respective programs (e.g., XBLAST and NBLAST) are used. See http://www.ncbi.nlm.nih.gov. A variant can also include, e.g., an internal deletion or insertion, a conservative or non-conservative substitution, or a combination of these variations from the sequence presented.

[00035] Soluble fragments are of great interest as these can competitively inhibit anthrax toxin binding to the ATR and thereby can protect cells from AT intoxication *in vivo* and *in vitro*. A fragment is soluble if it is not membrane-bound and is soluble in an aqueous fluid. The extracellular ATR domain is a soluble fragment of the ATR, as are fragments of that domain. Even though the VWA domain is formally identified as extending from amino acid 44 to 216 in the extracellular domain, more or fewer natively adjacent amino acids can be included in the fragment without compromising solubility or PA-binding. For example, a PA-binding fragment having the sequence of SEQ ID NO:2 beginning at any amino acid in the range from 27 to 43 and ending at any amino acid in the range from 221 to 321. A preferred soluble, PA-binding fragment extends from amino acid 42 to 222. Another preferred soluble PA-binding fragment includes a fragment of the ATR from amino acid 27 through amino acid 321. Likewise, any polypeptide fragment of these preferred fragments that retains PA-binding activity is within the scope of the invention. ATR in soluble form is effective in a monomeric form, as well as in multimeric forms such as dimeric, tetrameric, pentameric and higher oligomeric forms.

[00036] PA-binding polypeptides can include, therefore, SEQ ID NO:2, SEQ ID NO:6, SEQ ID NO:8, SEQ ID NO:10, a PA-binding fragment of SEQ ID NO:2, a PA-binding fragment of SEQ ID NO:6, a PA-binding fragment of SEQ ID NO:8, a PA-binding fragment of SEQ ID NO:10, a PA-binding polypeptide at least 80% identical to any of the foregoing fragments. The PA-binding polypeptides can also be provided as fusion proteins comprising any of the foregoing that can comprise still other non-natively adjacent amino acids for detecting, visualizing, isolating, or stabilizing the polypeptide. For example, PA binds to a soluble fusion protein of a hexahistidine tag, a T7 tag, and amino acids 41-227 of ATR.

[00037] Likewise, isolated polynucleotides having an uninterrupted nucleic acid sequence that encodes the aforementioned polypeptides and polypeptide fragments are also us ful in the

invention. The sequences that encode soluble, PA-binding polypeptide fragments of ATR are immediately apparent to the skilled artisan from the description of the relevant portions of the polypeptides, *supra*. An isolated nucleic acid containing the complement of any such polynucleotide is also within the scope of the present invention, as are polynucleotide and oligonucleotide fragments for use as molecular probes. The polynucleotides of the invention cannot encode SEQ ID NO:6, SEQ ID NO:8 or SEQ ID NO:10.

[00038] The present invention also relates to an isolated polynucleotide and its complement, without regard to source, where the polynucleotide hybridizes under stringent or moderately stringent hybridization conditions to SEQ ID NO:1, SEQ ID NO:5, SEQ ID 7, or SEQ ID NO:9 or to a fragment of any of the foregoing that encodes a soluble polypeptide that can bind to PA. As used herein, stringent conditions involve hybridizing at 68°C in 5x SSC/5x Denhardt's solution/1.0% SDS, and washing in 0.2x SSC/0.1% SDS +/- 100 μg/ml denatured salmon sperm DNA, at room temperature. Moderately stringent conditions include washing in the same buffer at 42°C. Additional guidance regarding such conditions is readily available in the art, for example, by Sambrook et al., 1989, Molecular Cloning, A Laboratory Manual, Cold Spring Harbor Press, N.Y.; and Ausubel et al. (eds.), 1995, Current Protocols in Molecular Biology, (John Wiley & Sons, N.Y.) at Unit 2.10.

[00039] In a related aspect, any polynucleotide of the invention can be provided in a vector in a manner known to those skilled in the art. The vector can be a cloning vector or an expression vector. In an expression vector, the polypeptide-encoding polynucleotide is under the transcriptional control of one or more non-native expression control sequences, such as a promoter not natively adjacent to the polynucleotide, such that the encoded polypeptide can be produced when the vector is delivered into a compatible host cell that supports expression of an polypeptide encoded on a vector, for example by electroporation or transfection, or transcribed and translated in a cell-free transcription and translation system. Such cell-based and cell-free systems are well known to the skilled artisan. Cells comprising an insert-containing vector of the invention are themselves within the scope of the present invention, without regard to whether the vector is extrachromosomal or integrated in the genome.

[00040] A skilled artisan in possession of the polypeptides and polynucleotides of the invention can also identify agents that can reduce or prevent the effect of AT on a host having on

the cell surface at least a portion of the ATR. The effect altered can relate, for example, to (1) susceptibility of the host cell to AT damage, (2) integration of ATR into the cell membrane, (3) binding between ATR and PA, (4) PA heptamerization, (5) uptake of PA and ATR complex into cells, and (6) the translocation of toxin into host cell cytoplasm. The method includes separately exposing a plurality of putative agents in the presence of AT to a plurality of cells, comparing the effect of AT on the cells in the presence and absence of the agent, and identifying at least one agent that alters an effect of AT on the cells.

[00041] The skilled artisan can readily evaluate the typical effects of AT and can observe variations in those effects in the presence of a putative altering agent. For example, susceptibility to AT damage can be evaluated by exposing host cells to AT. Integration of newly formed ATR into the host cell membrane can be evaluated by labeling newly synthesized proteins in the host cell and immunopreticipating ATR from the cellular membrane fraction of the host cell. Binding of wild-type ATR to PA can be evaluated with fluorescent labeled anti-PA antibody. PA heptamerization can be evaluated by several techniques including native polyacrylamide gel electrophoresis, gel filtration, and western blotting. Uptake of PA-ATR complex can be evaluated by binding PA to ATR at 4°C, increasing the temperature to 37°C to allow endocytosis, shifting the temperature back to 4°C, and incubating cells with fluorescent labeled anti-PA antibodies. Toxin translocation into the host cell cytoplasm can be evaluated as described in Wesche et al, 1998, which is incorporated herein by reference as if set forth in its entirety.

[00042] The agents screened can be, for example, dominant negative mutant ATRs (encoded by a mutant polynucleotide sequence, which can be provided in an expression vector), a high molecular weight molecule such as a polypeptide (including, e.g., a mutant AT, a soluble ATR, a mono- or polyclonal antibody to an ATR, to PA, or to an ATR/PA complex), a polysaccharide, a lipid, a nucleic acid, a low molecular weight organic or inorganic molecule, or the like. Antibodies can be produced by administering to a non-human animal an immunogenic, PA-binding fragment of a polypeptide which can be, e.g., SEQ ID NO:2, SEQ ID NO:6, SEQ ID NO:8, SEQ ID NO:10, a polypeptide at least 80% identical to any of the foregoing and a fusion protein comprising any of the foregoing, and then obtaining the desired antibodies using known methods.

[00043] Chemical libraries for screening putative agents, including peptide libraries, are

readily available to the skilled artisan. Examples include those from ASINEX (i.e. the Combined Wisdom Library of 24,000 manually synthesized organic molecules) and from CHEMBRIDGE CORPORATION (i.e. the DIVERSet™ library of 50,000 manually synthesized chemical compounds; the SCREEN-Set™ library of 24,000 manually synthesized chemical compounds; the CNS-Set™ library of 11,000 compounds; the Cherry-Pick™ library of up to 300,000 compounds) and linear library, multimeric library and cyclic library (Tecnogen (Italy)). Once an agent with desired activity is identified, a library of derivatives of that agent can be screened for better agents. Phage display is also a suitable approach for finding novel inhibitors of the interaction between PA and ATR.

[00044] Another aspect of the present invention relates to ATR ligands other than PA and methods for identifying ATR ligands. As ATR is expressed in many cell types, it likely has other natural ligands. To identify these other ligands, a polypeptide that contains an ATR VWA domain, preferably an entire extracellular domain can be provided in soluble or tethered form, e.g., in a chromatographic column. Preferably, the ectodomain of ATR can be provided as a fusion protein that also a contains rabbit IgG constant region, a GST domain or a hexahistidine tag. This fusion protein can be immobilized on a chromatographic column using known methods. A cell extract can be passed over the column. A ligand is identified when binding is observed between the ectodomain and a compound present in the cell extract. The identified ligand can be used in methods for identifying agents that alter an effect of AT, to identify an agent that selectively inhibits PA-ATR binding. It is also desirable to use the other ligands and the ATR in comparative high throughput screening methods for identifying small molecules that do not interfere with natural ligand binding to ATR, but which do prevent or reduce binding of ATR to anthrax toxin.

[00045] The present invention also relates to reducing cellular damage caused by AT, which can be achieved by administering an agent for reducing the ATR level, inhibiting the binding between ATR and AT, or by reducing downstream ATR activity after AT binding. For example, an antisense oligonucleotide can reduce or prevent expression of *atr* using delivery methods known to the skilled artisan, thus reducing the cellular ATR level. An ATR-anthrax binding inhibition agent can inhibit the binding between ATR and AT. Dominant negative ATRs can block downstream ATR activities required for AT toxicity. The agents used for reducing AT

damage to cells can be administered to a human or non-human animal, preferably in a standard pharmaceutical carrier, in an amount effective to reduce or eliminate anthrax toxicity.

[00046] A 20-25mer antisense oligonucleotide can be directed against 5' end of the *atr* message with phosphorothioate derivatives on the last three base pairs on the 3' end and the 5' end to enhance the half life and stability of the oligonucleotides. A carrier for an antisense oligonucleotide can be used. An example of a suitable carrier is cationic liposomes. For example, an oligonucleotide can be mixed with cationic liposomes prepared by mixing l-alpha dioleylphatidylcelthanolamine with dimethldioctadecylammonium bromide in a ratio of 5:2 in 1 ml of chloroform. The solvent will be evaporated and the lipids resuspended by sonication in 10 ml of saline. Another way to use an antisense oligonucleotide is to engineer it into a vector so that the vector can produce an antisense cRNA that blocks the translation of the mRNAs encoding for ATR. Similarly, RNAi techniques, which are now being applied to mammalian systems, are also suited for inhibiting ATR expression (see Zamore, Nat. Struct. Biol. 8:746:750 (2001), incorporated herein by reference as if set forth in its entirety).

[00047] The present invention also relates to a method for detecting *atr* mRNA or ATR protein in a sample. Such detection can be readily accomplished by using oligonucleotide or polynucleotide probes for *atr* mRNA, or antibodies for ATR protein. In a related aspect, the antibodies made and identified as being able to bind to ATR can also be used to separate ATR from a sample.

[00048] The present invention also relates to a cell line that does not contain ATR from a parent cell line that contains ATR, and methods for making same. The present invention provides that it is possible for cells lacking ATR to survive. In the example described below, a cell line that does not contain ATR was created using mutagenesis and screening. Now that the *atr* cDNA sequence is identified in the present invention, many other methods for generating a cell line that does not express *atr* become feasible, such as homologous recombination. In addition to these methods, the cell lines generated, including the one described in the example below, are themselves within the scope of the present invention.

[00049] The invention also provides molecules and methods for specifically targeting and killing cells of interest by delivering, e.g., AT or LF to the cell. Soluble ATR molecules can be coupled to a ligand or to a single chain antibody selected for targeting to the cell of interest (e.g.,

a ligand that binds a receptor presented on a tumor cell surface). The coupling is most readily accomplished by producing a fusion protein that encodes both the ATR binding portion and the ligand or single chain antibody molecule. The ligand or single chain antibody domains simply serve to attach the toxin to cells with the cognate surface markers. The toxin or factor is preloaded onto the ATR portion before exposing the coupled molecules to the targeted cells. This is similar in principle to the previously described for retroviral targeting using soluble retroviral receptor-ligand bridge proteins and retroviral receptor-single chain antibody bridge proteins. See Snitkovsky and Young, Proc. Natl. Acad. Sci. USA 95:7063-7068 (1998); Boerger et al. Proc. Natl. Acad. Sci. USA 96:9687-9872 (1999) and Snitkovsky et al., J. Virol. 74:9540-9545 (2000), and Snitkovsky et al., J. Virol. 75:1571-1575 (2001), each incorporated herein by reference as if set forth in its entirety.

[00050] The invention will be more fully understood upon consideration of the following non-limiting examples.

EXAMPLES

Methods

Mutagenesis and characterization of CHO-K1 cells

[00051] A mutant cell line lacking the receptor was generated, so that this defect could be genetically complemented. About 5 x 10^7 cells of the hypodiploid CHO-K1 cell line were treated at 37°C for 7 hr with medium containing 10 μg/ml ICR-191 (Sigma), a DNA alkylating agent that induces small deletions and frameshift mutations in genes, then washed twice. This treatment led to approximately 90% cell death.

[00052] The surviving mutagenized cells were then challenged with 8 μg/ml PA and 10 ng/ml LF_N-DTA, a fusion protein composed of the N-terminal 255 amino acids of LF linked to the catalytic A chain of diphtheria toxin. This recombinant toxin can kill CHO-K1 cells (in contrast to LF and PA) and it exploits the same LF/PA/receptor interactions that are required for the binding and entry of the native LF and EF proteins. After 4 days, surviving cells were replated and incubated for 3 days with medium containing PA and LF_N-DTA. Ten single-cell colonies (designated as CHO-R1.1 to CHO-R1.10) that survived toxin treatment were isolated 14 days later. In control experiments performed with non-mutagenized CHO-K1 cells, no toxin-resistant cell clones were detected.

[00053] One of the mutagenized clones (CHO-R1.1) was chosen for further analysis. CHO-R1.1 cells were found to be fully susceptible to killing by diphtheria toxin (DT) by measuring ^{3}H-leucine incorporation into cellular proteins after exposure to the toxin, thus ruling out the possibility that resistance to PA/LF_N-DTA was due to a defect in the pathway of DT action. To test directly whether CHO-R1.1 cells lacked the receptor, flow cytometric analysis was performed after the cells were incubated at 4°C for 2 hr in medium containing 40 to 80 nM PA-K563C coupled at mutated residue 563 to Oregon Green maleimide (Molecular Probes) ("OGPA"). The treated cells were washed twice with medium and analysed using a Becton Dickinson FACSCalibur flow cytometer. CHO-R1.1 cells were significantly impaired in their ability to bind to OGPA as compared to the parental cell line, suggesting that these mutagenized cells had lost expression of the putative PA receptor gene. Similar analysis of the other nine mutant CHO-R1 clones demonstrated that they were also defective in binding to OGPA.

cDNA complementation

[00054] In an attempt to complement the PA binding defect of CHO-R1.1 cells, the cells were transduced with a retrovirus-based cDNA library (Clontech) prepared from human HeLa cells that express the PA receptor. This cDNA library is contained in a murine leukemia virus (MLV) vector that is packaged into pseudotyped virus particles (MLV[VSV-G]) containing the broad host-range G protein of vesicular stomatitis virus (VSV-G). Retrovirus-based cDNA libraries are useful for genetic complementation approaches since they can deliver a limited number of stably expressed cDNA molecules per cell. These molecules can be rapidly re-isolated by PCR amplification using MLV vector-specific oligonucleotide primers.

[00055] Approximately 5 x 10^5 CHO-R1.1 cells were transduced with about 10^7 infectious units (complexity of library = 2 x 10^6 independent clones) of the pLIB-based cDNA library (Clontech; cat.# HL8002BB) produced in the 293GPG packaging cell line. Three days later, cells were incubated with medium containing 80 nM OGPA and the top 0.1% of fluorescent cells were then isolated by sorting using a Becton Dickinson FACSVantageSE instrument. Cells were sorted based on their binding of OGPA in combination with an anti-PA polyclonal serum and an allophycocyanin (APC) conjugated secondary antibody. To isolate those that contained the putative PA receptor cDNA clone, these cells were expanded and subjected to four additional rounds of sorting using OGPA as above, as well as a 1:500 dilution of a rabbit anti-PA polyclonal serum along with a 1:500 dilution of an APC-conjugated secondary antibody (Molecular probes). OGPA-single positive (round 2) or OGPA/APC-double positive (rounds 3-5) cells were recovered (the top 20%, 1%, 5%, and 50% of fluorescent cells for rounds 2, 3, 4, and 5 respectively) and expanded after each round of sorting.

[00056] This led to the isolation of a cell population in which greater than 90% of the cells bound OGPA. This complemented cell population contained at least seven unique cDNA inserts that were obtained by the PCR amplification method described above. Each cDNA was gel purified, subcloned back into the parent pLIB vector and packaged into MLV(VSV-G) virions so that it could be tested for its ability to complement the PA-binding defect of CHO-R1.1 cells. One cDNA clone of approximately 1.5 kb (designated as *ATR*) restored PA binding to CHO-R1.1 cells. This clone also dramatically enhanced the binding of PA to parental CHO-K1 cells.

[00057] Furthermore, the *ATR* cDNA clone fully restored LF_N-DTA/PA toxin sensitivity to CHO-R1.1 cells. In this test, CHO-R1.1 cells and CHO-K1 cells were either not transduced or

transduced with the MLV vector encoding ATR; these cells were treated with 10^{-9} M LF_N-DTA and various concentrations of PA; medium containing 1 μCi/mL ^{3}H-leucine was then added to cells for 1 hr, and the amount of ^{3}H-leucine incorporated into cellular proteins was determined by trichloroacetic acid precipitation and liquid scintillation counting.

CDNA characterization

[00058] cDNA inserts were recovered from these cells by PCR amplification of genomic DNA samples using oligonucleotide primers specific for the MLV vector according to the manufacturers instructions (Clontech). Each cDNA was subcloned between the *Not*I and *Sal*I restriction enzyme sites of pLIB and the resulting plasmids were co-transfected into 293 cells with MLV gag/pol and VSV-G expression plasmids pMD.old.gagpol and pMD.G. Resulting pseudotyped virus particles were used to infect CHO-R1.1 and CHO-K1 cells followed by OGPA staining and FACS analysis as above.

[00059] Sequencing of the *ATR* cDNA clone revealed a single long open reading frame, encoding a 368 amino acid protein. Fig. 1 shows sequence alignment of ATR (SEQ ID NO:2) with the von Willebrand factor A domain consensus sequence (SEQ ID NO:3; VWA-CON), the I domain of integrin α2 (SEQ ID NO:4; α2), and TEM8 (SEQ ID NO:6). The secondary structural elements are based on the crystal structure of the α2 I domain. Conserved amino acids are boxed and identical amino acids are indicated by shaded boxes. The putative signal sequence is underlined. The five residues that form the MIDAS motif are indicated with asterisks. The putative transmembrane domains of ATR and TEM8 are indicated with a shaded box. Potential N-linked glycosylation sites in ATR and TEM8 are indicated by hatched boxes. The alignment was made using the programs ClustalW and ESPript 1.9.

[00060] The ATR protein is predicted to have a 27 amino acid long signal peptide, a 293 amino acid long extracellular domain with three putative N-linked glycosylation sites, a 23 amino acid long putative transmembrane region, and a short cytoplasmic tail. A BLAST search revealed that the first 364 amino acids of ATR are identical to a protein encoded by the human *TEM8* cDNA clone (Genbank accession number NM 032208). The C-terminal ends of ATR and the TEM8 protein then diverge, presumably as a consequence of alternative splicing, such that ATR has a cytoplasmic tail of only 25 amino acids whereas TEM8 is predicted to have a 221 amino acid long cytoplasmic tail. The most notable feature of ATR is the presence of an

extracellular von Willebrand Factor type A (VWA) domain, located between residues 44 and 216.

[00061] The cytoplasmic tail of ATR contains an acidic cluster (AC motif) (EESEE) that is similar to a motif found in the cytoplasmic tail of furin which specifies basolateral sorting of this protease in polarized epithelial cells. This may be significant because the PA receptor localizes to the basolateral surface of polarized epithelial cells and it is expected that the receptor and the protease needed to bind and activate PA would be co-localized to allow for efficient entry of anthrax toxins.

Cloning and Expression of T7-ATR$_{41-227}$

[00062] A fusion protein having a hexahistidine tag, a T7 tag, and amino acids 41 to 227 of ATR (the I domain) was constructed, expressed and purified from *E. coli* cells as follows. A DNA fragment encoding amino acids 41-227 of ATR was cloned into the *Bam*H1 and *Eco*R1 sites of pET28A (Novagen) to generate pET28A-ATR$_{41-227}$. BL21 (DE3) cells (Stratagene) containing pET28A-ATR$_{41-227}$ were grown at 37°C to an OD_{600} of 0.6, induced with 1 mM isopropyl-β-D-thiogalactopyranoside for 4 hr and harvested by centrifugation. The cells from 1.5 L of culture were resuspended in 25 mL of 50 mM Tris-HCl pH 8.0, 2 mM dithiothreitol (DTT), 1 mM phenylmethylsulfonyl fluoride and were passed through a French press. One milligram of DNAse I (Roche) was added to the cell lysate, which was then sonicated for 1 min and centrifuged at 21,000 *g* for 20 min. The pellet was resuspended in 25 mL of 50 mM Tris-HCl pH 8.0, 2 mM DTT and centrifuged at 21,000 g for 20 min. This wash step was repeated once. T7-ATR$_{41-227}$ was solubilized and folded essentially as described previously.

[00063] When mixed with wild-type PA (on ice for 30 min), this construct was precipitated with polyclonal anti-PA serum (analyzed by SDS-PAGE and Western blot using anti-T7 antibody conjugated to horseradish peroxidase). The interaction between PA and T7-ATR$_{41-227}$ was impaired by the presence of EDTA (2 mM), demonstrating that the involvement of divalent cations in the interaction, and suggesting that the ATR MIDAS motif is involved in binding PA.

Interaction between PA and ATR

[00064] PA-N682S, a mutant form of PA isolated as described below and having an impaired ability to bind and intoxicate cells, did not bind to T7-ATR$_{41-227}$. The DNA encoding

Domain 4 of PA was mutagenized using error-prone PCR. Clones were expressed in *E. coli*, and lysates derived from these clones were added to CHO-K1 cells in combination with LF_N-DTA. Clones corresponding to lysates that did not kill CHO-K1 cells were sequenced and the N682S mutant clone was further characterized as having Ser in place of Asn at position 682.

[00065] PA-N682S was shown to have an impaired ability to bind cells as follows: CHO-K1 cells were incubated with $2x10^{-8}$ M trypsin-nicked PA (wild-type or N682S) for 1 hr, washed with PBS, resuspended in SDS sample buffer and run on a 4-20 % polyacrylamide SDS gel, and PA was visualized by Western blotting. In the experiment in which PA-N682S was shown to have an impaired ability to intoxicate cells, CHO-K1 cells were incubated with LF_N-DTA (10^{-9} M) and various concentrations of wild-type PA or PA-N682S mutant, and cell viability was determined.

[00066] To confirm that PA binds directly to ATR, co-immunoprecipitations (using a polyclonal serum specific for PA and protein A agarose) were performed with an extracellular fragment of ATR and either the wild-type or a receptor binding-deficient mutant form of PA. A mixture of 5 μg PA (WT or N682S) and 2 μg T7-ATR_{41-227} (in 20 mM Tris-HCl pH 8.0, 150 mM NaCl, 0.1 mg bovine serum albumin per mL) was incubated on ice for 30 min in the presence or absence of 2 mM EDTA. Anti-PA polyclonal serum (10 μL) was added to this solution and incubated on ice for an additional 1 hr. Protein A agarose (Santa Cruz Biotechnology) was added and the solution was rotated at 4°C for 1 hr, then washed four times with 20 mM Tris-HCl pH 8.0, 150 mM NaCl. Approximately one third of the mixture was subjected to SDS-PAGE, transferred to nitrocellulose and probed with anti-T7 antibody conjugated to horseradish peroxidase (Novagen).

[00067] In addition, a fusion protein containing GST and the PA receptor-binding domain (D4) (GST-D4) bound T7-ATR_{41-227}, while GST did not. DNA encoding amino acids 595 to 735 of PA (domain 4) was cloned into pGEX-4T-1 (Pharmacia Biotechnology). This vector encoded the GST-D4 fusion protein. GST-D4 was coupled to glutathione sepharose at a concentration of 4 mg GST-D4 per mL according to manufacturer's instructions (Pharmacia Biotechnology). GST or GST-D4 coupled to glutathione sepharose was mixed with 2 μg of T7-ATR_{41-227} and 250 μg of *E. coli* extract in a volume of 250 μL for 1 hr at 4°C. The beads were washed 4 times with 20 mM Tris-HCl pH 8.0, 150 mM NaCl. One half of the suspension was subjected to SDS-

PAGE, transferred to nitrocellulose, and probed with anti-T7 antibody coupled to horseradish peroxidase.

[00068] Taken together, the experiments described above demonstrate a direct and specific interaction between the VWA/I domain of ATR and the receptor-binding domain of PA. Given this direct interaction, we reasoned that ATR_{41-227} might protect CHO-K1 cells from killing by PA and LF_N-DTA. This idea was tested by incubating (37°C for 4 hr) CHO-K1 cells with an increasing amount of T7-ATR_{41-227} in the presence of a constant amount of PA (10^{-10} M)/LF_N-DTA (2.5×10^{-11} M), and then measuring the subsequent effect on protein synthesis. T7-ATR_{41-227} was an effective inhibitor of toxin action, inhibiting toxin activity by 50% and 100% at concentrations of 80 nM and 500 nM respectively. T7-ATR_{41-227} did not, however, inhibit diphtheria toxin.

[00069] The present invention is not intended to be limited to the foregoing, but encompasses all such modifications and variations as come within the scope of the appended claims.

CLAIMS

WE CLAIM:

1. A isolated polypeptide selected from the group consisting of SEQ ID NO:2, a PA-binding fragment of SEQ ID NO:2, a PA-binding fragment of SEQ ID NO:6, a PA-binding fragment of SEQ ID NO:8, a PA-binding fragment of SEQ ID NO:10, a PA-binding polypeptide at least 80% identical to any of the foregoing fragments, and a fusion protein comprising any of the foregoing.

2. The isolated polypeptide of claim 1 wherein the polypeptide is a soluble polypeptide.

3. The isolated polypeptide of claim 1, wherein the PA-binding fragment of SEQ ID NO:2 begins at any amino acid in the range from 27 to 43 and ends at any amino acid in the range from 221 to 321.

4. The isolated polypeptide of claim 1 having an amino acid sequence set forth in SEQ ID NO:2.

5. An isolated polynucleotide or complement thereof, the polynucleotide encoding a polypeptide selected from the group consisting of SEQ ID NO:2, a PA-binding fragment of SEQ ID NO:2, a PA-binding fragment of SEQ ID NO:6, a PA-binding fragment of SEQ ID NO:8, a PA-binding fragment of SEQ ID NO:10, a PA-binding polypeptide at least 80% identical to any of the foregoing fragments, and a fusion protein comprising any of the foregoing, the polynucleotide being unable to encode SEQ ID NO:6, SEQ ID NO:8 or SEQ ID NO:10.

6. The isolated polynucleotide of claim 5, wherein the PA-binding fragment of SEQ ID NO:2 begins at any amino acid in the range from 27 to 43 and ends at any amino acid in the range from 221 to 321.

7. The isolated polynucleotide of claim 5 comprising SEQ ID NO:1 from position 104 to 1207 or the complement thereof.

8. The isolated polynucleotide of claim 5 comprising SEQ ID NO:1 or the complement thereof.

9. The isolated polynucleotide of claim 5, wherein the polynucleotide encodes a soluble polypeptide.

10. An isolated polynucleotide or complement thereof, the polynucleotide hybridizing under stringent or moderately stringent hybridization conditions to all or a portion of SEQ ID NO:1 and encoding a soluble, PA-binding polypeptide.

11. A vector comprising a polynucleotide selected from the group consisting of a polynucleotide of claim 5 and a polynucleotide of claim 10.

12. The vector of claim 11, further comprising a non-native expression control sequence operably linked to the polynucleotide.

13. A host cell comprising a vector of claim 11.

14. A method for making an antibody, the method comprising the step of: administering to a non-human animal an immunogenic, PA-binding fragment of a polypeptide selected from the group consisting of SEQ ID NO:2, SEQ ID NO:6, SEQ ID NO:8, SEQ ID NO:10, a polypeptide

at least 80% identical to any of the foregoing and a fusion protein comprising any of the foregoing.

15. A method for identifying an agent that inhibits binding between protective antigen (PA) and anthrax toxin receptor, the method comprising the steps of:

combining protective antigen (PA) and a polypeptide selected from the group consisting of SEQ ID NO:2, SEQ ID NO:6, SEQ ID NO:8, SEQ ID NO:10, a PA-binding fragment of SEQ ID NO:2, a PA-binding fragment of SEQ ID NO:6, a PA-binding fragment of SEQ ID NO:8, a PA-binding fragment of SEQ ID NO:10, a PA-binding polypeptide at least 80% identical to any of the foregoing, and a fusion protein comprising any of the foregoing, separately with and without a putative binding-inhibiting agent;

comparing binding between PA and the polypeptide with and without the putative agent; and

identifying a decrease in binding with the putative agent, the decrease being an indication that the test agent inhibits the binding of PA to the anthrax toxin receptor.

16. A method for treating anthrax in a human or non-human animal, the method comprising the step of:

administering to the animal an agent that inhibits binding between protective antigen (PA) and anthrax toxin receptor at a level effective to reduce the severity of anthrax.

17. A method as claimed in claim 16, wherein the agent that inhibits binding between PA and the anthrax toxin receptor is selected from the group consisting of SEQ ID NO:2, SEQ ID NO:6, SEQ ID NO:8, SEQ ID NO:10, a PA-binding fragment of SEQ ID NO:2, a PA-binding fragment of SEQ ID NO:6, a PA-binding fragment of SEQ ID NO:8, a PA-binding fragment of SEQ ID NO:10, a PA-binding polypeptide at least 80% identical to any of the foregoing, a fusion protein comprising any of the foregoing, a monoclonal antibody, a polyclonal antibody, a polysaccharide, a lipid, and a nucleic acid.

18. A cultured cell having a cell membrane having an exterior surface, the exterior surface displaying no receptor for anthrax toxin protective antigen.

19. A method for producing an anthrax toxin receptor, the method including the step of:

transcribing a polynucleotide that encodes an anthrax toxin receptor operably linked to an upstream expression control sequence, the receptor being selected from the group consisting of SEQ ID NO:2, SEQ ID NO:6, SEQ ID NO:8, SEQ ID NO:10, a PA-binding fragment of SEQ ID NO:2, a PA-binding fragment of SEQ ID NO:6, a PA-binding fragment of SEQ ID NO:8, a PA-binding fragment of SEQ ID NO:10, a PA-binding polypeptide at least 80% identical to any of the foregoing, and a fusion protein comprising any of the foregoing, to produce an mRNA; and

translating the mRNA to produce the anthrax toxin receptor.

20. A method as claimed in Claim 19, wherein the polynucleotide is operably linked to the expression control sequence in an expression vector, and wherein the expression vector is delivered into a host cell, the expression control sequence being operable in the host cell.

21. A method as claimed in Claim 19, wherein at least one of the transcribing and translating steps are performed *in vitro*.

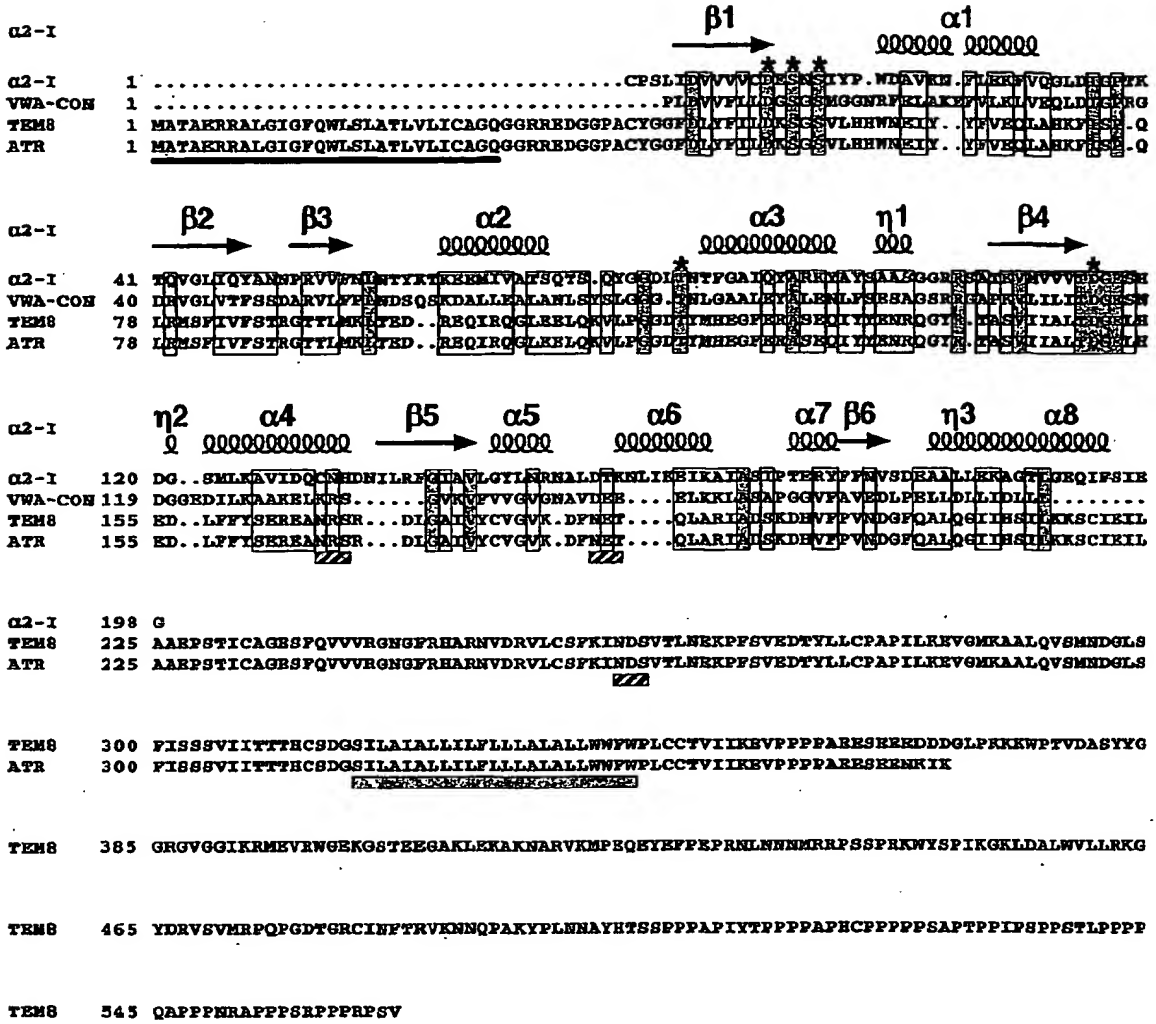
α2-I
β1
α1
α2-I 1 .....CPSLIDVVVVCDESNSIYP.WDAVKN.FLKKFVQGLDIGPTK
VWA-CON 1 .....PLDVVFLLDGSGSMGGNRFKLAKEFVLKIVEQLDVGPDR
TEM8 1 MATAERRALGIGFQWLSLATLVLICAGQGGRREDGGPACYGGFDLYFDLDNSGSVLHHWNEIY..YFVEQLAHKFFSN-Q
ATR 1 MATAERRALGIGFQWLSLATLVLICAGQGGRREDGGPACYGGFDLYFDLDNSGSVLHHWNEIY..YFVEQLAHKFFSN-Q
α2-I
β2
β3
α2
α3
η1
β4
α2-I 41 TQVGLIQYANSPRVVVPMSTNTYRTKEEMIVATSQTS.QYGGDLTNTFGAIQYARKYAYSAASGGRRSATKVMVVVTDGES
VWA-CON 40 DRVGLVTFSSDARVLFTLNDSQSKDALLEALANLSYSLGSA.NLGAALTTALEMLFSESAGSRIGAPKVLIIITDGES
TEM8 78 LRMSFIVFSTRGTTLMKLTED..RSQIRQGLEELQKVLPSGLDMIMHEGFEKRSEQIYDSNRQGIK.MASVIIALESLH
ATR 78 LRMSFIVFSTRGTTLMKLTED..RSQIRQGLEELQKVLPSGLDMIMHEGFEKRSEQIYDSNRQGIK.TASVIIALESLH
α2-I
η2
α4
β5
α5
α6
α7
β6
η3
α8
α2-I 120 DG..SMLKAVIDQCNHDNILRFGIAVLGYLNRNALDTKNLIKSIKAIASIPTERYFFNVSDEAALLEKAGTLGEQIFSIE
VWA-CON 119 DGGRDILKAAKELKRS.....GVKVFVVGVGNAVDEE....ELKKLASAPGGVFAVEDLPELLDLLIDILE.....
TEM8 155 ED..LFFYSERRANRSR...DLGAIVYCVGVK.DFNET....QLARIASKDHVFFVNDGFQALQGIIHSILKKSCIEIL
ATR 155 ED..LFFYSERRANRSR...DLGAIVYCVGVK.DFNET....QLARIASKDHVFFVNDGFQALQGIIHSILKKSCIEIL
α2-I 198 G
TEM8 225 AARPSTICAGESFQVVVRGNGFRHARNVDRVLCSFKINDSVTLNEKPFSVEDTYLLCPAPILKEVGMKAALQVSMNDGLS
ATR 225 AARPSTICAGESFQVVVRGNGFRHARNVDRVLCSFKINDSVTLNEKPFSVEDTYLLCPAPILKEVGMKAALQVSMNDGLS
TEM8 300 FISSSVIITTTHCSDGSILAIALLILFLLLALALLWNFWPLCCTVIIKEVPPPPARESREEDDDGLPKKWPTVDASYYG
ATR 300 FISSSVIITTTHCSDGSILAIALLILFLLLALALLWNFWPLCCTVIIKEVPPPPARESREENKIK
TEM8 385 GRGVGGIKRMEVRWGEKGSTEEGAKLEKAKNARVKMPEQEYEFPEPRNLNNNMRRPSSPRKWYSPIKGKLDALWVLLRKG
TEM8 465 YDRVSVMRPQPGDTGRCINFTRVKNNQPAKYPLNNAYHTSSPPPAPIYTPPPPAPHCPPPPPSAPTPPIPSPPSTLPPPP
TEM8 545 QAPPPNRAPPPSRPPPRPSV

FIG 1

SEQUENCE LISTING

<110> Young, John A.T.
Bradley, Kenneth A.
Collier, Robert J.
Mogridge, Jeremy S.

<120> Anthrax Toxin Receptor

<130> 960296.97745

<140>
<141>

<150> 60/251,481
<151> 2000-12-05

<160> 10

<170> PatentIn Ver. 2.1

<210> 1
<211> 1414
<212> DNA
<213> Homo sapiens

<220>
<221> CDS
<222> (104)..(1207)

<400> 1
aggacccgcg aggaagggcc cgcggatggc gcgtccctga gggtcgtggc gagttcgcgg 60

agcgtgggaa ggagcggacc ctgctctccc cgggctgcgg gcc atg gcc acg gcg 115
Met Ala Thr Ala
1

gag cgg aga gcc ctc ggc atc ggc ttc cag tgg ctc tct ttg gcc act 163
Glu Arg Arg Ala Leu Gly Ile Gly Phe Gln Trp Leu Ser Leu Ala Thr
5 10 15 20

ctg gtg ctc atc tgc gcc ggg caa ggg gga cgc agg gag gat ggg ggt 211
Leu Val Leu Ile Cys Ala Gly Gln Gly Gly Arg Arg Glu Asp Gly Gly
25 30 35

cca gcc tgc tac ggc gga ttt gac ctg tac ttc att ttg gac aaa tca 259
Pro Ala Cys Tyr Gly Gly Phe Asp Leu Tyr Phe Ile Leu Asp Lys Ser
40 45 50

gga agt gtg ctg cac cac tgg aat gaa atc tat tac ttt gtg gaa cag 307
Gly Ser Val Leu His His Trp Asn Glu Ile Tyr Tyr Phe Val Glu Gln
55 60 65

ttg gct cac aaa ttc atc agc cca cag ttg aga atg tcc ttt att gtt 355
Leu Ala His Lys Phe Ile Ser Pro Gln Leu Arg Met Ser Phe Ile Val
70 75 80

```
ttc tcc acc cga gga aca acc tta atg aaa ctg aca gaa gac aga gaa   403
Phe Ser Thr Arg Gly Thr Thr Leu Met Lys Leu Thr Glu Asp Arg Glu
 85                  90                  95                 100

caa atc cgt caa ggc cta gaa gaa ctc cag aaa gtt ctg cca gga gga   451
Gln Ile Arg Gln Gly Leu Glu Glu Leu Gln Lys Val Leu Pro Gly Gly
                105                 110                 115

gac act tac atg cat gaa gga ttt gaa agg gcc agt gag cag att tat   499
Asp Thr Tyr Met His Glu Gly Phe Glu Arg Ala Ser Glu Gln Ile Tyr
            120                 125                 130

tat gaa aac aga caa ggg tac agg aca gcc agc gtc atc att gct ttg   547
Tyr Glu Asn Arg Gln Gly Tyr Arg Thr Ala Ser Val Ile Ile Ala Leu
        135                 140                 145

act gat gga gaa ctc cat gaa gat ctc ttt ttc tat tca gag agg gag   595
Thr Asp Gly Glu Leu His Glu Asp Leu Phe Phe Tyr Ser Glu Arg Glu
    150                 155                 160

gct aat agg tct cga gat ctt ggt gca att gtt tac tgt gtt ggt gtg   643
Ala Asn Arg Ser Arg Asp Leu Gly Ala Ile Val Tyr Cys Val Gly Val
165                 170                 175                 180

aaa gat ttc aat gag aca cag ctg gcc cgg att gcg gac agt aag gat   691
Lys Asp Phe Asn Glu Thr Gln Leu Ala Arg Ile Ala Asp Ser Lys Asp
                185                 190                 195

cat gtg ttt ccc gtg aat gac ggc ttt cag gct ctg caa ggc atc atc   739
His Val Phe Pro Val Asn Asp Gly Phe Gln Ala Leu Gln Gly Ile Ile
            200                 205                 210

cac tca att ttg aag aag tcc tgc atc gaa att cta gca gct gaa cca   787
His Ser Ile Leu Lys Lys Ser Cys Ile Glu Ile Leu Ala Ala Glu Pro
        215                 220                 225

tcc acc ata tgt gca gga gag tca ttt caa gtt gtc gtg aga gga aac   835
Ser Thr Ile Cys Ala Gly Glu Ser Phe Gln Val Val Val Arg Gly Asn
    230                 235                 240

ggc ttc cga cat gcc cgc aac gtg gac agg gtc ctc tgc agc ttc aag   883
Gly Phe Arg His Ala Arg Asn Val Asp Arg Val Leu Cys Ser Phe Lys
245                 250                 255                 260

atc aat gac tcg gtc aca ctc aat gag aag ccc ttt tct gtg gaa gac   931
Ile Asn Asp Ser Val Thr Leu Asn Glu Lys Pro Phe Ser Val Glu Asp
                265                 270                 275

act tat tta ctg tgt cca gcg cct atc tta aaa gaa gtt ggc atg aaa   979
Thr Tyr Leu Leu Cys Pro Ala Pro Ile Leu Lys Glu Val Gly Met Lys
            280                 285                 290

gct gca ctc cag gtc agc atg aac gat ggc ctc tct ttt atc tcc agt  1027
Ala Ala Leu Gln Val Ser Met Asn Asp Gly Leu Ser Phe Ile Ser Ser
        295                 300                 305
```

```
tct gtc atc atc acc acc aca cac tgt tct gac ggt tcc atc ctg gcc   1075
Ser Val Ile Ile Thr Thr Thr His Cys Ser Asp Gly Ser Ile Leu Ala
    310                 315                 320

atc gcc ctg ctg atc ctg ttc ctg ctc cta gcc ctg gct ctc ctc tgg   1123
Ile Ala Leu Leu Ile Leu Phe Leu Leu Leu Ala Leu Ala Leu Leu Trp
325                 330                 335                 340

tgg ttc tgg ccc ctc tgc tgc act gtg att atc aag gag gtc cct cca   1171
Trp Phe Trp Pro Leu Cys Cys Thr Val Ile Ile Lys Glu Val Pro Pro
                345                 350                 355

ccc cct gcc gag gag agt gag gaa aat aaa ata aaa taacaagaag        1217
Pro Pro Ala Glu Glu Ser Glu Glu Asn Lys Ile Lys
            360                 365

aagaaagaaa gaaatcccac agaaacagat aacctaacac agcccgtgca acgtatttta 1277

tacaatgctc tgaaaatcat agtctcaatc tagacagtct tttcctctag ttccctgtat 1337

tcaaatccca gtgtctaaca ttcaataaat agctatatga aatcaaaaaa aaaaaaaaaa 1397

aaaaaaaaaa aaaaaaa                                               1414
```

<210> 2
<211> 368
<212> PRT
<213> Homo sapiens

<400> 2

```
Met Ala Thr Ala Glu Arg Arg Ala Leu Gly Ile Gly Phe Gln Trp Leu
  1               5                  10                  15

Ser Leu Ala Thr Leu Val Leu Ile Cys Ala Gly Gln Gly Gly Arg Arg
             20                  25                  30

Glu Asp Gly Gly Pro Ala Cys Tyr Gly Gly Phe Asp Leu Tyr Phe Ile
         35                  40                  45

Leu Asp Lys Ser Gly Ser Val Leu His His Trp Asn Glu Ile Tyr Tyr
     50                  55                  60

Phe Val Glu Gln Leu Ala His Lys Phe Ile Ser Pro Gln Leu Arg Met
 65                  70                  75                  80

Ser Phe Ile Val Phe Ser Thr Arg Gly Thr Thr Leu Met Lys Leu Thr
                 85                  90                  95

Glu Asp Arg Glu Gln Ile Arg Gln Gly Leu Glu Glu Leu Gln Lys Val
            100                 105                 110

Leu Pro Gly Gly Asp Thr Tyr Met His Glu Gly Phe Glu Arg Ala Ser
        115                 120                 125

Glu Gln Ile Tyr Tyr Glu Asn Arg Gln Gly Tyr Arg Thr Ala Ser Val
    130                 135                 140
```

```
Ile Ile Ala Leu Thr Asp Gly Glu Leu His Glu Asp Leu Phe Phe Tyr
145                 150                 155                 160

Ser Glu Arg Glu Ala Asn Arg Ser Arg Asp Leu Gly Ala Ile Val Tyr
                165                 170                 175

Cys Val Gly Val Lys Asp Phe Asn Glu Thr Gln Leu Ala Arg Ile Ala
            180                 185                 190

Asp Ser Lys Asp His Val Phe Pro Val Asn Asp Gly Phe Gln Ala Leu
        195                 200                 205

Gln Gly Ile Ile His Ser Ile Leu Lys Lys Ser Cys Ile Glu Ile Leu
    210                 215                 220

Ala Ala Glu Pro Ser Thr Ile Cys Ala Gly Glu Ser Phe Gln Val Val
225                 230                 235                 240

Val Arg Gly Asn Gly Phe Arg His Ala Arg Asn Val Asp Arg Val Leu
                245                 250                 255

Cys Ser Phe Lys Ile Asn Asp Ser Val Thr Leu Asn Glu Lys Pro Phe
            260                 265                 270

Ser Val Glu Asp Thr Tyr Leu Leu Cys Pro Ala Pro Ile Leu Lys Glu
        275                 280                 285

Val Gly Met Lys Ala Ala Leu Gln Val Ser Met Asn Asp Gly Leu Ser
    290                 295                 300

Phe Ile Ser Ser Ser Val Ile Ile Thr Thr Thr His Cys Ser Asp Gly
305                 310                 315                 320

Ser Ile Leu Ala Ile Ala Leu Leu Ile Leu Phe Leu Leu Leu Ala Leu
                325                 330                 335

Ala Leu Leu Trp Trp Phe Trp Pro Leu Cys Cys Thr Val Ile Ile Lys
            340                 345                 350

Glu Val Pro Pro Pro Pro Ala Glu Glu Ser Glu Glu Asn Lys Ile Lys
        355                 360                 365
```

<210> 3
<211> 180
<212> PRT
<213> Artificial Sequence

<220>
<223> Description of Artificial Sequence:von Willebrand factor A domain consensus sequence

<400> 3

```
Pro Leu Asp Val Val Phe Leu Leu Asp Gly Ser Gly Ser Met Gly Gly
  1               5                  10                  15
```

```
Asn Arg Phe Glu Leu Ala Lys Glu Phe Val Leu Lys Leu Val Glu Gln
            20                  25                  30

Leu Asp Ile Gly Pro Arg Gly Asp Arg Val Gly Leu Val Thr Phe Ser
        35                  40                  45

Ser Asp Ala Arg Val Leu Phe Pro Leu Asn Asp Ser Gln Ser Lys Asp
    50                  55                  60

Ala Leu Leu Glu Ala Leu Ala Asn Leu Ser Tyr Ser Leu Gly Gly Gly
65                  70                  75                  80

Thr Asn Leu Gly Ala Ala Leu Glu Tyr Ala Leu Glu Asn Leu Phe Ser
                85                  90                  95

Glu Ser Ala Gly Ser Arg Arg Gly Ala Pro Lys Val Leu Ile Leu Ile
            100                 105                 110

Thr Asp Gly Glu Ser Asn Asp Gly Gly Glu Asp Ile Leu Lys Ala Ala
        115                 120                 125

Lys Glu Leu Lys Arg Ser Gly Val Lys Val Phe Val Val Gly Val Gly
    130                 135                 140

Asn Ala Val Asp Glu Glu Glu Leu Lys Lys Leu Ala Ser Ala Pro Gly
145                 150                 155                 160

Gly Val Phe Ala Val Glu Asp Leu Pro Glu Leu Leu Asp Leu Leu Ile
                165                 170                 175

Asp Leu Leu Leu
            180
```

<210> 4
<211> 198
<212> PRT
<213> Homo sapiens

<400> 4

```
Cys Pro Ser Leu Ile Asp Val Val Val Val Cys Asp Glu Ser Asn Ser
  1               5                  10                  15

Ile Tyr Pro Trp Asp Ala Val Lys Asn Phe Leu Glu Lys Phe Val Gln
            20                  25                  30

Gly Leu Asp Ile Gly Pro Thr Lys Thr Gln Val Gly Leu Ile Gln Tyr
        35                  40                  45

Ala Asn Asn Pro Arg Val Val Phe Asn Leu Asn Thr Tyr Lys Thr Lys
    50                  55                  60

Glu Glu Met Ile Val Ala Thr Ser Gln Thr Ser Gln Tyr Gly Gly Asp
65                  70                  75                  80
```

```
Leu Thr Asn Thr Phe Gly Ala Ile Gln Tyr Ala Arg Lys Tyr Ala Tyr
                 85                  90                  95

Ser Ala Ser Gly Gly Arg Arg Ser Ala Ala Thr Lys Val Met Val Val
            100                 105                 110

Val Thr Asp Gly Glu Ser His Asp Gly Ser Met Leu Lys Ala Val Ile
        115                 120                 125

Asp Gln Cys Asn His Asp Asn Ile Leu Arg Phe Gly Ile Ala Val Leu
    130                 135                 140

Gly Tyr Leu Asn Arg Asn Ala Leu Asp Thr Lys Asn Leu Ile Lys Glu
145                 150                 155                 160

Ile Lys Ala Ile Ala Ser Ile Pro Thr Glu Arg Tyr Phe Phe Asn Val
                165                 170                 175

Ser Asp Glu Ala Ala Leu Leu Glu Lys Ala Gly Thr Leu Gly Glu Gln
            180                 185                 190

Ile Phe Ser Ile Glu Gly
        195


<210> 5
<211> 5540
<212> DNA
<213> Homo sapiens

<220>
<221> CDS
<222> (144)..(1835)

<400> 5
aattgcttcc ggggagttgc gagggagcga gggggaataa aggacccgcg aggaagggcc 60

cgcggatggc gcgtccctga gggtcgtggc gagttcgcgg agcgtgggaa ggagcggacc 120

ctgctctccc cgggctgcgg gcc atg gcc acg gcg gag cgg aga gcc ctc ggc 173
                        Met Ala Thr Ala Glu Arg Arg Ala Leu Gly
                          1               5                  10

atc ggc ttc cag tgg ctc tct ttg gcc act ctg gtg ctc atc tgc gcc   221
Ile Gly Phe Gln Trp Leu Ser Leu Ala Thr Leu Val Leu Ile Cys Ala
                 15                  20                  25

ggg caa ggg gga cgc agg gag gat ggg ggt cca gcc tgc tac ggc gga   269
Gly Gln Gly Gly Arg Arg Glu Asp Gly Gly Pro Ala Cys Tyr Gly Gly
             30                  35                  40

ttt gac ctg tac ttc att ttg gac aaa tca gga agt gtg ctg cac cac   317
Phe Asp Leu Tyr Phe Ile Leu Asp Lys Ser Gly Ser Val Leu His His
         45                  50                  55
```

```
tgg aat gaa atc tat tac ttt gtg gaa cag ttg gct cac aaa ttc atc   365
Trp Asn Glu Ile Tyr Tyr Phe Val Glu Gln Leu Ala His Lys Phe Ile
     60                  65                  70

agc cca cag ttg aga atg tcc ttt att gtt ttc tcc acc cga gga aca   413
Ser Pro Gln Leu Arg Met Ser Phe Ile Val Phe Ser Thr Arg Gly Thr
 75                  80                  85                  90

acc tta atg aaa ctg aca gaa gac aga gaa caa atc cgt caa ggc cta   461
Thr Leu Met Lys Leu Thr Glu Asp Arg Glu Gln Ile Arg Gln Gly Leu
                 95                 100                 105

gaa gaa ctc cag aaa gtt ctg cca gga gga gac act tac atg cat gaa   509
Glu Glu Leu Gln Lys Val Leu Pro Gly Gly Asp Thr Tyr Met His Glu
            110                 115                 120

gga ttt gaa agg gcc agt gag cag att tat tat gaa aac aga caa ggg   557
Gly Phe Glu Arg Ala Ser Glu Gln Ile Tyr Tyr Glu Asn Arg Gln Gly
        125                 130                 135

tac agg aca gcc agc gtc atc att gct ttg act gat gga gaa ctc cat   605
Tyr Arg Thr Ala Ser Val Ile Ile Ala Leu Thr Asp Gly Glu Leu His
    140                 145                 150

gaa gat ctc ttt ttc tat tca gag agg gag gct aat agg tct cga gat   653
Glu Asp Leu Phe Phe Tyr Ser Glu Arg Glu Ala Asn Arg Ser Arg Asp
155                 160                 165                 170

ctt ggt gca att gtt tac tgt gtt ggt gtg aaa gat ttc aat gag aca   701
Leu Gly Ala Ile Val Tyr Cys Val Gly Val Lys Asp Phe Asn Glu Thr
                175                 180                 185

cag ctg gcc cgg att gcg gac agt aag gat cat gtg ttt ccc gtg aat   749
Gln Leu Ala Arg Ile Ala Asp Ser Lys Asp His Val Phe Pro Val Asn
            190                 195                 200

gac ggc ttt cag gct ctg caa ggc atc atc cac tca att ttg aag aag   797
Asp Gly Phe Gln Ala Leu Gln Gly Ile Ile His Ser Ile Leu Lys Lys
        205                 210                 215

tcc tgc atc gaa att cta gca gct gaa cca tcc acc ata tgt gca gga   845
Ser Cys Ile Glu Ile Leu Ala Ala Glu Pro Ser Thr Ile Cys Ala Gly
    220                 225                 230

gag tca ttt caa gtt gtc gtg aga gga aac ggc ttc cga cat gcc cgc   893
Glu Ser Phe Gln Val Val Val Arg Gly Asn Gly Phe Arg His Ala Arg
235                 240                 245                 250

aac gtg gac agg gtc ctc tgc agc ttc aag atc aat gac tcg gtc aca   941
Asn Val Asp Arg Val Leu Cys Ser Phe Lys Ile Asn Asp Ser Val Thr
                255                 260                 265

ctc aat gag aag ccc ttt tct gtg gaa gat act tat tta ctg tgt cca   989
Leu Asn Glu Lys Pro Phe Ser Val Glu Asp Thr Tyr Leu Leu Cys Pro
            270                 275                 280
```

```
gcg cct atc tta aaa gaa gtt ggc atg aaa gct gca ctc cag gtc agc   1037
Ala Pro Ile Leu Lys Glu Val Gly Met Lys Ala Ala Leu Gln Val Ser
        285                 290                 295

atg aac gat ggc ctc tct ttt atc tcc agt tct gtc atc atc acc acc   1085
Met Asn Asp Gly Leu Ser Phe Ile Ser Ser Ser Val Ile Ile Thr Thr
    300                 305                 310

aca cac tgt tct gac ggt tcc atc ctg gcc atc gcc ctg ctg atc ctg   1133
Thr His Cys Ser Asp Gly Ser Ile Leu Ala Ile Ala Leu Leu Ile Leu
315                 320                 325                 330

ttc ctg ctc cta gcc ctg gct ctc ctc tgg tgg ttc tgg ccc ctc tgc   1181
Phe Leu Leu Leu Ala Leu Ala Leu Leu Trp Trp Phe Trp Pro Leu Cys
                335                 340                 345

tgc act gtg att atc aag gag gtc cct cca ccc cct gcc gag gag agt   1229
Cys Thr Val Ile Ile Lys Glu Val Pro Pro Pro Pro Ala Glu Glu Ser
            350                 355                 360

gag gaa gaa gat gat gat ggt ctg cct aag aaa aag tgg cca acg gta   1277
Glu Glu Glu Asp Asp Asp Gly Leu Pro Lys Lys Lys Trp Pro Thr Val
        365                 370                 375

gac gcc tct tat tat ggt ggg aga ggc gtt gga ggc att aaa aga atg   1325
Asp Ala Ser Tyr Tyr Gly Gly Arg Gly Val Gly Gly Ile Lys Arg Met
    380                 385                 390

gag gtt cgt tgg gga gaa aag ggc tcc aca gaa gaa ggt gct aag ttg   1373
Glu Val Arg Trp Gly Glu Lys Gly Ser Thr Glu Glu Gly Ala Lys Leu
395                 400                 405                 410

gaa aag gca aag aat gca aga gtc aag atg ccg gag cag gaa tat gaa   1421
Glu Lys Ala Lys Asn Ala Arg Val Lys Met Pro Glu Gln Glu Tyr Glu
                415                 420                 425

ttc cct gag ccg cga aat ctc aac aac aat atg cgt cgg cct tct tcc   1469
Phe Pro Glu Pro Arg Asn Leu Asn Asn Asn Met Arg Arg Pro Ser Ser
            430                 435                 440

ccc cgg aag tgg tac tct cca atc aag gga aaa ctc gat gcc ttg tgg   1517
Pro Arg Lys Trp Tyr Ser Pro Ile Lys Gly Lys Leu Asp Ala Leu Trp
        445                 450                 455

gtc cta ctg agg aaa gga tat gat cgt gtg tct gtg atg cgt cca cag   1565
Val Leu Leu Arg Lys Gly Tyr Asp Arg Val Ser Val Met Arg Pro Gln
    460                 465                 470

cca gga gac acg ggg cgc tgc atc aac ttc acc agg gtc aag aac aac   1613
Pro Gly Asp Thr Gly Arg Cys Ile Asn Phe Thr Arg Val Lys Asn Asn
475                 480                 485                 490

cag cca gcc aag tac cca ctc aac aac gcc tac cac acc tcc tcg ccg   1661
Gln Pro Ala Lys Tyr Pro Leu Asn Asn Ala Tyr His Thr Ser Ser Pro
                495                 500                 505
```

```
cct cct gcc ccc atc tac act ccc cca cct cct gcg ccc cac tgc cct   1709
Pro Pro Ala Pro Ile Tyr Thr Pro Pro Pro Pro Ala Pro His Cys Pro
            510                 515                 520

ccc ccg ccc ccc agc gcc cct acc cct ccc atc ccg tcc cca cct tcc   1757
Pro Pro Pro Pro Ser Ala Pro Thr Pro Pro Ile Pro Ser Pro Pro Ser
        525                 530                 535

acc ctt ccc cct cct ccc cag gct cca cct ccc aac agg gca cct cct   1805
Thr Leu Pro Pro Pro Pro Gln Ala Pro Pro Pro Asn Arg Ala Pro Pro
    540                 545                 550

ccc tcc cgc cct cct cca agg cct tct gtc tagagcccaa agttcctgct     1855
Pro Ser Arg Pro Pro Pro Arg Pro Ser Val
555                 560

ctgggctctc tcagaaactt caggagatgt tagaacaagt ctttccagtt agagaagagg 1915

agtggtgata aagcccactg accttcacac attctaaaaa ttggttggca atgccagtat 1975

accaacaatc atgatcagct gaaagaaaca gatattttaa attgccagaa aacaaatgat 2035

gaggcaacta cagtcagatt tatagccagc catctatcac ctctagaagg ttccagagac 2095

agtgaaactg caagatgctc tcaacaggat tatgtctcat ggagaccagt aagaaaatca 2155

tttatctgaa ggtgaaatgc agagttggat aagaaataca ttgctgggtt tctaaaatgc 2215

tgccttcctg cctctactcc acctccatcc ctggactttg gacccttggc ctaggagcct 2275

aaggaccttc acccctgtgc accacccaag aaagaggaaa actttgccta caactttgga 2335

aatgctgggg tccctggtgt ggtaagaaac tcaacatcag acgggtatgc agaaggatgt 2395

tcttctggga tttgcaggta cataaaaaat gtatggcatc ttttccttgc aaattcttcc 2455

agtttccaag tgagaagggg agcaggtgtt tactgatgga aaaggtatgt tgctatgttg 2515

atgtgtaagt gaaatcagtt gtgtgcaata gacaggggcg tattcatggg agcatcagcc 2575

agtttctaaa acccacaggc catcagcagc tagaggtggc tggctttggc cagacatgga 2635

ccctaaatca acagacaatg gcattgtcga agagcaacct gttaatgaat catgttaaaa 2695

atcaaggttt ggcttcagtt taaatcactt gaggtatgaa gtttatcctg ttttccagag 2755

ataaacataa gttgatcttc ccaaaatacc atcattagga cctatcacac aatatcacta 2815

gtttttttg tttgtttgtt ttttgttttt tttcttggta aagccatgca ccacagactt 2875

ctgggcagag ctgagagaca atggtcctga cataataagg atctttgatt aacccccata 2935

aggcatgtgt gtgtatacaa atatacttct ctttggcttt tcgacataga acctcagctg 2995

ttaaccaagg ggaaatacat cagatctgca acacagaaat gctctgcctg aaatttccac 3055

catgcctagg actcacccca tttatccagg tctttctgga tctgtttaat caataagccc 3115
```

tataatcact tgctaaacac tgggcttcat cacccaggga taaaaacaga gatcattgtc 3175

ttggacctcc tgcatcagcc tattcaaaat tatctctctc tctagctttc cacaaatcct 3235

aaaattcctg tcccaagcca cccaaattct cagatctttt ctggaacaag gcagaatata 3295

aaataaatat acatttagtg gcttgggcta tggtctccaa agatccttca aaaatacatc 3355

aagccagctt cattcactca ctttacttag aacagagata taagggcctg ggatgcattt 3415

attttatcaa taccaatttt tgtggccatg gcagacattg ctaatcaatc acagcactat 3475

ttcctattaa gcccactgat ttcttcacaa tccttctcaa attacaattc caaagagccg 3535

ccactcaaca gtcagatgaa cccaacagtc agatgagaga aatgaaccct acttgctatc 3595

tctatcttag aaagcaaaaa caaacaggag tttccaggga gaatgggaaa gccaggggge 3655

ataaaaggta cagtcagggg aaaatagatc taggcagagt gccttagtca gggaccacgg 3715

gcgctgaatc tgcagtgcca acaccaaact gacacatctc caggtgtacc tccaacccta 3775

gccttctccc acagctgcct acaacagagt ctcccagcct tctcagagag ctaaaaccag 3835

aaatttccag actcatgaaa gcaacccccc agcctctccc caaccctgcc gcattgtcta 3895

atttttagaa cactaggctt cttctttcat gtagttcctc ataagcaggg gccagaatat 3955

ctcagccacc tgcagtgaca ttgctggacc cctgaaaacc attccatagg agaatgggtt 4015

ccccaggctc acagtgtaga gacattgagc ccatcacaac tgttttgact gctggcagtc 4075

taaaacagtc cacccacccc atggcactgc cgcgtgattc ccgcggccat tcagaagttc 4135

aagccgagat gctgacgttg ctgagcaacg agatggtgag catcagtgca aatgcaccat 4195

tcagcacatc agtcatatgc ccagtgcagt tacaagatgt tgtttcggca aagcattttg 4255

atggaatagg gaactgcaaa tgtatgatga ttttgaaaag gctcagcagg atttgttctt 4315

aaaccgactc agtgtgtcat ccccggttat ttagaattac agttaagaag gagaaacttc 4375

tataagactg tatgaacaag gtgatatctt catagtgggc tattacaggc aggaaaatgt 4435

tttaactggt ttacaaaatc catcaatact tgtgtcattc cctgtaaaag gcaggagaca 4495

tgtgattatg atcaggaaac tgcacaaaat tattgttttc agcccccgtg ttattgtcct 4555

tttgaactgt tttttttta ttaaagccaa atttgtgttg tatatattcg tattccatgt 4615

gttagatgga agcatttcct atccagtgtg aataaaaaga acagttgtag taaattatta 4675

taaagccgat gatatttcat ggcaggttat tctaccaagc tgtgcttgtt ggtttttccc 4735

atgactgtat tgcttttata aatgtacaaa tagttactga aatgacgaga cccttgtttg 4795

cacagcatta ataagaacct tgataagaac catattctgt tgacagccag ctcacagttt 4855

cttgcctgaa gcttggtgca ccctccagtg agacacaaga tctctctttt accaaagttg 4915

agaacagagc tggtggatta attaatagtc ttcgatatct ggccatgggt aacctcattg 4975

taactatcat cagaatgggc agagatgatc ttgaagtgtc acatacacta aagtccaaac 5035

actatgtcag atgggggtaa aatccattaa agaacaggaa aaaataatta taagatgata 5095

agcaaatgtt tcagcccaat gtcaacccag ttaaaaaaaa aattaatgct gtgtaaaatg 5155

gttgaattag tttgcaaact atataaagac atatgcagta aaaagtctgt taatgcacat 5215

cctgtgggaa tggagtgttc taaccaattg ccttttcttg ttatctgagc tctcctatat 5275

tatcatactc agataaccaa attaaaagaa ttagaatatg atttttaata cacttaacat 5335

taaactcttc taactttctt ctttctgtga taattcagaa gatagttatg gatcttcaat 5395

gcctctgagt cattgttata aaaaatcagt tatcactata ccatgctata ggagactggg 5455

caaaacctgt acaatgacaa ccctggaagt tgctttttt aaaaaaataa taaatttctt 5515

aaatcaaaaa aaaaaaaaaa aaaaa 5540

<210> 6
<211> 564
<212> PRT
<213> Homo sapiens

<400> 6

```
Met Ala Thr Ala Glu Arg Arg Ala Leu Gly Ile Gly Phe Gln Trp Leu
  1               5                  10                  15

Ser Leu Ala Thr Leu Val Leu Ile Cys Ala Gly Gln Gly Gly Arg Arg
             20                  25                  30

Glu Asp Gly Gly Pro Ala Cys Tyr Gly Gly Phe Asp Leu Tyr Phe Ile
         35                  40                  45

Leu Asp Lys Ser Gly Ser Val Leu His His Trp Asn Glu Ile Tyr Tyr
     50                  55                  60

Phe Val Glu Gln Leu Ala His Lys Phe Ile Ser Pro Gln Leu Arg Met
 65                  70                  75                  80

Ser Phe Ile Val Phe Ser Thr Arg Gly Thr Thr Leu Met Lys Leu Thr
                 85                  90                  95

Glu Asp Arg Glu Gln Ile Arg Gln Gly Leu Glu Glu Leu Gln Lys Val
            100                 105                 110

Leu Pro Gly Gly Asp Thr Tyr Met His Glu Gly Phe Glu Arg Ala Ser
        115                 120                 125
```

```
Glu Gln Ile Tyr Tyr Glu Asn Arg Gln Gly Tyr Arg Thr Ala Ser Val
    130                 135                 140

Ile Ile Ala Leu Thr Asp Gly Glu Leu His Glu Asp Leu Phe Phe Tyr
145                 150                 155                 160

Ser Glu Arg Glu Ala Asn Arg Ser Arg Asp Leu Gly Ala Ile Val Tyr
                165                 170                 175

Cys Val Gly Val Lys Asp Phe Asn Glu Thr Gln Leu Ala Arg Ile Ala
            180                 185                 190

Asp Ser Lys Asp His Val Phe Pro Val Asn Asp Gly Phe Gln Ala Leu
        195                 200                 205

Gln Gly Ile Ile His Ser Ile Leu Lys Lys Ser Cys Ile Glu Ile Leu
    210                 215                 220

Ala Ala Glu Pro Ser Thr Ile Cys Ala Gly Glu Ser Phe Gln Val Val
225                 230                 235                 240

Val Arg Gly Asn Gly Phe Arg His Ala Arg Asn Val Asp Arg Val Leu
                245                 250                 255

Cys Ser Phe Lys Ile Asn Asp Ser Val Thr Leu Asn Glu Lys Pro Phe
            260                 265                 270

Ser Val Glu Asp Thr Tyr Leu Leu Cys Pro Ala Pro Ile Leu Lys Glu
        275                 280                 285

Val Gly Met Lys Ala Ala Leu Gln Val Ser Met Asn Asp Gly Leu Ser
    290                 295                 300

Phe Ile Ser Ser Ser Val Ile Ile Thr Thr Thr His Cys Ser Asp Gly
305                 310                 315                 320

Ser Ile Leu Ala Ile Ala Leu Leu Ile Leu Phe Leu Leu Leu Ala Leu
                325                 330                 335

Ala Leu Leu Trp Trp Phe Trp Pro Leu Cys Cys Thr Val Ile Ile Lys
            340                 345                 350

Glu Val Pro Pro Pro Pro Ala Glu Glu Ser Glu Glu Glu Asp Asp Asp
        355                 360                 365

Gly Leu Pro Lys Lys Lys Trp Pro Thr Val Asp Ala Ser Tyr Tyr Gly
    370                 375                 380

Gly Arg Gly Val Gly Gly Ile Lys Arg Met Glu Val Arg Trp Gly Glu
385                 390                 395                 400

Lys Gly Ser Thr Glu Glu Gly Ala Lys Leu Glu Lys Ala Lys Asn Ala
                405                 410                 415

Arg Val Lys Met Pro Glu Gln Glu Tyr Glu Phe Pro Glu Pro Arg Asn
            420                 425                 430
```

```
Leu Asn Asn Asn Met Arg Arg Pro Ser Ser Pro Arg Lys Trp Tyr Ser
        435                 440                 445

Pro Ile Lys Gly Lys Leu Asp Ala Leu Trp Val Leu Leu Arg Lys Gly
    450                 455                 460

Tyr Asp Arg Val Ser Val Met Arg Pro Gln Pro Gly Asp Thr Gly Arg
465                 470                 475                 480

Cys Ile Asn Phe Thr Arg Val Lys Asn Asn Gln Pro Ala Lys Tyr Pro
                485                 490                 495

Leu Asn Asn Ala Tyr His Thr Ser Ser Pro Pro Pro Ala Pro Ile Tyr
            500                 505                 510

Thr Pro Pro Pro Pro Ala Pro His Cys Pro Pro Pro Pro Pro Ser Ala
        515                 520                 525

Pro Thr Pro Pro Ile Pro Ser Pro Pro Ser Thr Leu Pro Pro Pro Pro
    530                 535                 540

Gln Ala Pro Pro Pro Asn Arg Ala Pro Pro Pro Ser Arg Pro Pro Pro
545                 550                 555                 560

Arg Pro Ser Val
```

<210> 7
<211> 2112
<212> DNA
<213> Homo sapiens

<220>
<221> CDS
<222> (113)..(1111)

<400> 7
ggggaataaa ggacccgcga ggaagggccc gcggatggcg cgtccctgag ggtcgtggcg 60

agttcgcgga gcgtgggaag gagcggaccc tgctctcccc gggctgcggg cc atg gcc 118
                                                        Met Ala
                                                          1

acg gcg gag cgg aga gcc ctc ggc atc ggc ttc cag tgg ctc tct ttg 166
Thr Ala Glu Arg Arg Ala Leu Gly Ile Gly Phe Gln Trp Leu Ser Leu
          5   10   15

gcc act ctg gtg ctc atc tgc gcc ggg caa ggg gga cgc agg gag gat 214
Ala Thr Leu Val Leu Ile Cys Ala Gly Gln Gly Gly Arg Arg Glu Asp
     20   25   30

ggg ggt cca gcc tgc tac ggc gga ttt gac ctg tac ttc att ttg gac 262
Gly Gly Pro Ala Cys Tyr Gly Gly Phe Asp Leu Tyr Phe Ile Leu Asp
 35   40   45   50

```
aaa tca gga agt gtg ctg cac cac tgg aat gaa atc tat tac ttt gtg   310
Lys Ser Gly Ser Val Leu His His Trp Asn Glu Ile Tyr Tyr Phe Val
                 55                  60                  65

gaa cag ttg gct cac aaa ttc atc agc cca cag ttg aga atg tcc ttt   358
Glu Gln Leu Ala His Lys Phe Ile Ser Pro Gln Leu Arg Met Ser Phe
            70                  75                  80

att gtt ttc tcc acc cga gga aca acc tta atg aaa ctg aca gaa gac   406
Ile Val Phe Ser Thr Arg Gly Thr Thr Leu Met Lys Leu Thr Glu Asp
        85                  90                  95

aga gaa caa atc cgt caa ggc cta gaa gaa ctc cag aaa gtt ctg cca   454
Arg Glu Gln Ile Arg Gln Gly Leu Glu Glu Leu Gln Lys Val Leu Pro
    100                 105                 110

gga gga gac act tac atg cat gaa gga ttt gaa agg gcc agt gag cag   502
Gly Gly Asp Thr Tyr Met His Glu Gly Phe Glu Arg Ala Ser Glu Gln
115                 120                 125                 130

att tat tat gaa aac aga caa ggg tac agg aca gcc agc gtc atc att   550
Ile Tyr Tyr Glu Asn Arg Gln Gly Tyr Arg Thr Ala Ser Val Ile Ile
                135                 140                 145

gct ttg act gat gga gaa ctc cat gaa gat ctc ttt ttc tat tca gag   598
Ala Leu Thr Asp Gly Glu Leu His Glu Asp Leu Phe Phe Tyr Ser Glu
            150                 155                 160

agg gag gct aat agg tct cga gat ctt ggt gca att gtt tac tgt gtt   646
Arg Glu Ala Asn Arg Ser Arg Asp Leu Gly Ala Ile Val Tyr Cys Val
        165                 170                 175

ggt gtg aaa gat ttc aat gag aca cag ctg gcc cgg att gcg gac agt   694
Gly Val Lys Asp Phe Asn Glu Thr Gln Leu Ala Arg Ile Ala Asp Ser
    180                 185                 190

aag gat cat gtg ttt ccc gtg aat gac ggc ttt cag gct ctg caa ggc   742
Lys Asp His Val Phe Pro Val Asn Asp Gly Phe Gln Ala Leu Gln Gly
195                 200                 205                 210

atc atc cac tca att ttg aag aag tcc tgc atc gaa att cta gca gct   790
Ile Ile His Ser Ile Leu Lys Lys Ser Cys Ile Glu Ile Leu Ala Ala
                215                 220                 225

gaa cca tcc acc ata tgt gca gga gag tca ttt caa gtt gtc gtg aga   838
Glu Pro Ser Thr Ile Cys Ala Gly Glu Ser Phe Gln Val Val Val Arg
            230                 235                 240

gga aac ggc ttc cga cat gcc cgc aac gtg gac agg gtc ctc tgc agc   886
Gly Asn Gly Phe Arg His Ala Arg Asn Val Asp Arg Val Leu Cys Ser
        245                 250                 255

ttc aag atc aat gac tcg gtc aca ctc aat gag aag ccc ttt tct gtg   934
Phe Lys Ile Asn Asp Ser Val Thr Leu Asn Glu Lys Pro Phe Ser Val
    260                 265                 270
```

```
gaa gat act tat tta ctg tgt cca gcg cct atc tta aaa gaa gtt ggc   982
Glu Asp Thr Tyr Leu Leu Cys Pro Ala Pro Ile Leu Lys Glu Val Gly
275                 280                 285                 290

atg aaa gct gca ctc cag gtc agc atg aac gat ggc ctc tct ttt atc   1030
Met Lys Ala Ala Leu Gln Val Ser Met Asn Asp Gly Leu Ser Phe Ile
                295                 300                 305

tcc agt tct gtc atc atc acc acc aca cac tgt agc ctc cac aaa att   1078
Ser Ser Ser Val Ile Ile Thr Thr His Cys Ser Leu His Lys Ile
            310                 315                 320

gca tca ggc ccc aca aca gct gct tgc atg gaa tagcagagaa taccgcctgc 1131
Ala Ser Gly Pro Thr Thr Ala Ala Cys Met Glu
        325                 330

tccctccgga cagcacactc ctgaaaacgg ggagagagga gccaaacatg ctcggtttac 1191

actttcctta tttactgaat gagtggaggg cagagacagg cctggagtta cgcacactga 1251

gtgccccaac atggaaagaa acatcaggag ggacaggaaa cgttccctcc ttaaccaaca 1311

gttttcaaga ccttactgga ggcactttat tggctacata atcactccat gcggtgggca 1371

tcaggcagaa tcctggtgca gacccaactt tgaggtggag gatttcacag tttctttatt 1431

ttgaacttcc cccaggctcc cactaattcc tctccattct atcctcctcc ctttcccaca 1491

aaagaaaaca gaaaggagca gcagtgtttg ataccgtatc atccagaggc ctggttctct 1551

cccattatag ggcaaacaag ccctggcaag atatttcact cccgccccat gccatgcatt 1611

aaaaatccaa aattgcctat attccacctg ccaagcaaga gatgctttca ttattgaagt 1671

tccaaatgta taccttttgag aacagtgcct tctcgtctta aaagagaggt cctcattttg 1731

tgagttggga gcagagggaa ttaaagaaag ccatgatgca gggatttggc cattcaagcc 1791

gggcagcctt cagagaatgt catccctaat gacacatgcc cgaatgaagg agcggggctg 1851

agcttgtcct gccttcgtat tgaatgttgc ctgtctgcct ccttaatagc gggcctctgt 1911

gtgagcattt gacaagactt aaaactattc attgaagaaa atggatgatc ccccaacagg 1971

aagatgcaac cccatgggct gcctgcttga ccacagaagt gcttccagct ccagttgctc 2031

atctgagaac tccccccacc acttgctgtt aaaattgtta aaattaaagg ccatgttgat 2091

tgaaaaaaaa aaaaaaaaaa a                                          2112
```

<210> 8
<211> 333
<212> PRT
<213> Homo sapiens

<400> 8

```
Met Ala Thr Ala Glu Arg Arg Ala Leu Gly Ile Gly Phe Gln Trp Leu
 1               5                  10                  15

Ser Leu Ala Thr Leu Val Leu Ile Cys Ala Gly Gln Gly Gly Arg Arg
            20                  25                  30

Glu Asp Gly Gly Pro Ala Cys Tyr Gly Gly Phe Asp Leu Tyr Phe Ile
        35                  40                  45

Leu Asp Lys Ser Gly Ser Val Leu His His Trp Asn Glu Ile Tyr Tyr
    50                  55                  60

Phe Val Glu Gln Leu Ala His Lys Phe Ile Ser Pro Gln Leu Arg Met
65                  70                  75                  80

Ser Phe Ile Val Phe Ser Thr Arg Gly Thr Thr Leu Met Lys Leu Thr
                85                  90                  95

Glu Asp Arg Glu Gln Ile Arg Gln Gly Leu Glu Glu Leu Gln Lys Val
            100                 105                 110

Leu Pro Gly Gly Asp Thr Tyr Met His Glu Gly Phe Glu Arg Ala Ser
        115                 120                 125

Glu Gln Ile Tyr Tyr Glu Asn Arg Gln Gly Tyr Arg Thr Ala Ser Val
    130                 135                 140

Ile Ile Ala Leu Thr Asp Gly Glu Leu His Glu Asp Leu Phe Phe Tyr
145                 150                 155                 160

Ser Glu Arg Glu Ala Asn Arg Ser Arg Asp Leu Gly Ala Ile Val Tyr
                165                 170                 175

Cys Val Gly Val Lys Asp Phe Asn Glu Thr Gln Leu Ala Arg Ile Ala
            180                 185                 190

Asp Ser Lys Asp His Val Phe Pro Val Asn Asp Gly Phe Gln Ala Leu
        195                 200                 205

Gln Gly Ile Ile His Ser Ile Leu Lys Lys Ser Cys Ile Glu Ile Leu
    210                 215                 220

Ala Ala Glu Pro Ser Thr Ile Cys Ala Gly Glu Ser Phe Gln Val Val
225                 230                 235                 240

Val Arg Gly Asn Gly Phe Arg His Ala Arg Asn Val Asp Arg Val Leu
                245                 250                 255

Cys Ser Phe Lys Ile Asn Asp Ser Val Thr Leu Asn Glu Lys Pro Phe
            260                 265                 270

Ser Val Glu Asp Thr Tyr Leu Leu Cys Pro Ala Pro Ile Leu Lys Glu
        275                 280                 285

Val Gly Met Lys Ala Ala Leu Gln Val Ser Met Asn Asp Gly Leu Ser
    290                 295                 300
```

```
Phe Ile Ser Ser Ser Val Ile Ile Thr Thr Thr His Cys Ser Leu His
305                 310                 315                 320

Lys Ile Ala Ser Gly Pro Thr Thr Ala Ala Cys Met Glu
            325                 330
```

<210> 9
<211> 1436
<212> DNA
<213> Homo sapiens

<220>
<221> CDS
<222> (380)..(1033)

<400> 9

```
aattgcttcc ggggagttgc gagggagcga gggggaataa aggacccgcg aggaagggcc 60

cgcggatggc gcgtccctga gggtcgtggc gagttcgcgg agcgtgggaa ggagcggacc 120

ctgctctccc cgggctgcgg gccatggcca cggcggagcg gagagccctc ggcatcggct 180

tccagtggct ctcacggcca ctctggtgct catctgcgcc gggcaagggg gacgcaggga 240

ggatgggggt ccagcctgct acggcggatt tgacctgtac ttcattttgg acaaatcagg 300

aagtgtgctg caccactgga atgaaatcta ttactttgtg gaacagttgg ctcacaaatt 360

catcagccca cagttgaga atg tcc ttt att gtt ttc tcc acc cga gga aca  412
                    Met Ser Phe Ile Val Phe Ser Thr Arg Gly Thr
                      1               5                  10

acc tta atg aaa ctg aca gaa gac aga gaa caa atc cgt caa ggc cta  460
Thr Leu Met Lys Leu Thr Glu Asp Arg Glu Gln Ile Arg Gln Gly Leu
            15                  20                  25

gaa gaa ctc cag aaa gtt ctg cca gga gga gac act tac atg cat gaa  508
Glu Glu Leu Gln Lys Val Leu Pro Gly Gly Asp Thr Tyr Met His Glu
        30                  35                  40

gga ttt gaa agg gcc agt gag cag att tat tat gaa aac aga caa ggg  556
Gly Phe Glu Arg Ala Ser Glu Gln Ile Tyr Tyr Glu Asn Arg Gln Gly
    45                  50                  55

tac agg aca gct agc gtc atc att gct ttg act gat gga gaa ctc cat  604
Tyr Arg Thr Ala Ser Val Ile Ile Ala Leu Thr Asp Gly Glu Leu His
60                  65                  70                  75

gaa gat ctc ttt ttc tat tca gag agg gag gct aat agg tct cga gat  652
Glu Asp Leu Phe Phe Tyr Ser Glu Arg Glu Ala Asn Arg Ser Arg Asp
                80                  85                  90

ctt ggt gca att gtt tac tgt gtt ggt gtg aaa gat ttc aat gag aca  700
Leu Gly Ala Ile Val Tyr Cys Val Gly Val Lys Asp Phe Asn Glu Thr
            95                  100                 105
```

```
cag ctg gcc cgg att gcg gac agt aag gat cat gtg ttt ccc gtg aat   748
Gln Leu Ala Arg Ile Ala Asp Ser Lys Asp His Val Phe Pro Val Asn
        110                 115                 120

gac ggc ttt cag gct ctg caa ggc atc atc cac tca att ttg aag aag   796
Asp Gly Phe Gln Ala Leu Gln Gly Ile Ile His Ser Ile Leu Lys Lys
    125                 130                 135

tcc tgc atc gaa att cta gca gct gaa cca tcc acc ata tgt gca gga   844
Ser Cys Ile Glu Ile Leu Ala Ala Glu Pro Ser Thr Ile Cys Ala Gly
140                 145                 150                 155

gag tca ttt caa gtt gtc gtg aga gga aac ggc ttc cga cat gcc cgc   892
Glu Ser Phe Gln Val Val Val Arg Gly Asn Gly Phe Arg His Ala Arg
                160                 165                 170

aac gtg gac agg gtc ctc tgc agc ttc aag atc aat gac tcg gtc aca   940
Asn Val Asp Arg Val Leu Cys Ser Phe Lys Ile Asn Asp Ser Val Thr
            175                 180                 185

ctc agt aag tcc ttg cag agt cca tgg gtt tct tcg aca agt ggc ttc   988
Leu Ser Lys Ser Leu Gln Ser Pro Trp Val Ser Ser Thr Ser Gly Phe
        190                 195                 200

aag gaa ggg aat tcc cac cct tgt ctt cca gca agg cca cac aca       1033
Lys Glu Gly Asn Ser His Pro Cys Leu Pro Ala Arg Pro His Thr
    205                 210                 215

tgaaaccagc agaaaagagt cttatttgct ggaaagaccc ccagcaaggg catagtgagc 1093

ccttacagtg gttccagtca gaaaaggcac cacttgggtg ggcacagccc catgggtgtc 1153

caacttggta agcagagcaa ggctggactt gagtccccgt cctccacaaa acacagagcc 1213

acaagcccca gccctgcagc agccctccgg aagcagcggg gcactggttt ccttgtcccc 1273

tgccatctac cgagtggctc actctcaggt gggagtgctg gtgatggtta attaggactg 1333

cagaaacatg agcctcctta acaaagtatt gggactctta agggtaagtg tgaaaaagga 1393

atggtctaaa tgcattaatc ttgaataaac cgaaaaccaa acc                   1436
```

<210> 10
<211> 218
<212> PRT
<213> Homo sapiens

<400> 10

```
Met Ser Phe Ile Val Phe Ser Thr Arg Gly Thr Thr Leu Met Lys Leu
  1               5                  10                  15

Thr Glu Asp Arg Glu Gln Ile Arg Gln Gly Leu Glu Glu Leu Gln Lys
             20                  25                  30
```

```
Val Leu Pro Gly Gly Asp Thr Tyr Met His Glu Gly Phe Glu Arg Ala
         35                  40                  45

Ser Glu Gln Ile Tyr Tyr Glu Asn Arg Gln Gly Tyr Arg Thr Ala Ser
     50                  55                  60

Val Ile Ile Ala Leu Thr Asp Gly Glu Leu His Glu Asp Leu Phe Phe
 65                  70                  75                  80

Tyr Ser Glu Arg Glu Ala Asn Arg Ser Arg Asp Leu Gly Ala Ile Val
                85                  90                  95

Tyr Cys Val Gly Val Lys Asp Phe Asn Glu Thr Gln Leu Ala Arg Ile
            100                 105                 110

Ala Asp Ser Lys Asp His Val Phe Pro Val Asn Asp Gly Phe Gln Ala
        115                 120                 125

Leu Gln Gly Ile Ile His Ser Ile Leu Lys Lys Ser Cys Ile Glu Ile
    130                 135                 140

Leu Ala Ala Glu Pro Ser Thr Ile Cys Ala Gly Glu Ser Phe Gln Val
145                 150                 155                 160

Val Val Arg Gly Asn Gly Phe Arg His Ala Arg Asn Val Asp Arg Val
                165                 170                 175

Leu Cys Ser Phe Lys Ile Asn Asp Ser Val Thr Leu Ser Lys Ser Leu
            180                 185                 190

Gln Ser Pro Trp Val Ser Ser Thr Ser Gly Phe Lys Glu Gly Asn Ser
        195                 200                 205

His Pro Cys Leu Pro Ala Arg Pro His Thr
    210                 215
```